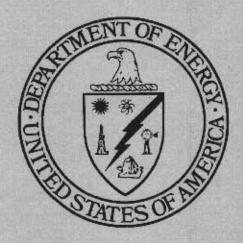


#### Sandia National Laboratories / New Mexico

#### PROPOSAL FOR NO FURTHER ACTION ENVIRONMENTAL RESTORATION PROJECT SITE 50, OLD CENTRIFUGE SITE OPERABLE UNIT 1309

**June 1995** 

Environmental Restoration Project

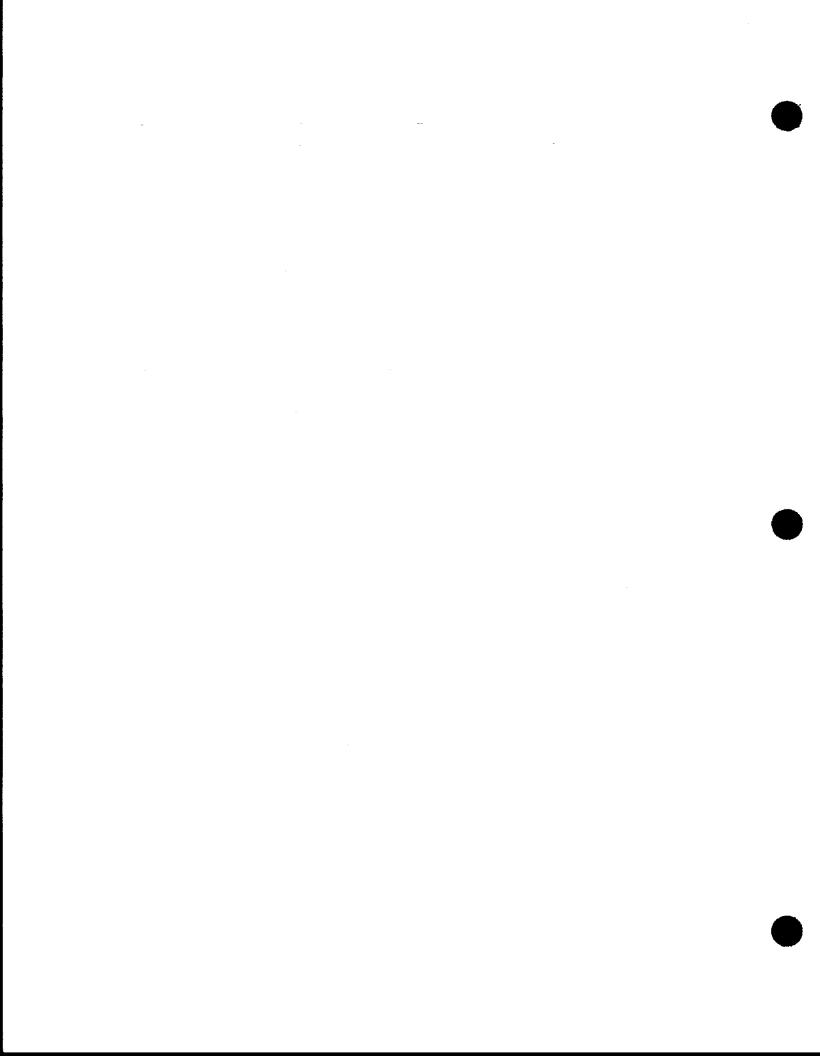


United States Department of Energy Albuquerque Operations Office

# PROPOSAL FOR NO FURTHER ACTION

Site 50, Old Centrifuge Site Operable Unit 1309

SANDIA NATIONAL LABORATORIES/NEW MEXICO



#### 1. Introduction

#### 1.1 ER Site Identification Number and Name

Sandia National Laboratories/New Mexico (SNL/NM) is proposing a risk-based no further action (NFA) decision for Environmental Restoration (ER) Site 50, Old Centrifuge Site, Operable Unit (OU) 1309. ER Site 50 is listed in the Hazardous and Solid Waste Amendment (HSWA) Module IV (EPA August 1993) of the SNL/NM Resource Conservation and Recovery Act (RCRA) Hazardous Waste Management Facility Permit (NM5890110518) (EPA August 1992).

#### 1.2 SNL/NM Risk Based NFA Process

This proposal for a determination of a risked-based NFA decision has been prepared using the criteria presented in Section 4.5.3 of the SNL/NM Program Implementation Plan (PIP) (SNL/NM February 1994). Specifically, this proposal will "contain information demonstrating that this Solid Wase Management Unit (SWMU) has never contained constituents of concern that may pose a threat to human health or the environment" [as proposed in the Code of Federal Regulations (CFR), Section 40 Part 264.51(a) (2)] (EPA July 1990). The HSWA Module IV contains the same requirements for an NFA demonstration:

Based on the results of the RFI [RCRA Facility Investigation] and other relevant information, the Permittee may submit an application to the Administrative Authority for a Class III permit modification under 40 CFR 270.42 (c) to terminate the RFI/CMS [corrective measures study] process for a specific unit. This permit modification application must contain information demonstrating that there are no releases of hazardous waste including hazardous constituents from a particular SWMU at the facility that pose threats to human health and/or the environment, as well as additional information required in 40 CFR 270.42 (c) (EPA August 1993).

For a risk-based proposal, a SWMU is eligible for an NFA determination if the NFA criterion established by the SNL/NM permit is met. This criterion, found in Section M.1 of the permit, is as follows: "[T]here are no releases of hazardous waste including hazardous constituents...that pose threats to human health and/or the environment..." This risk-base proposal contains information needed to make the NFA determination.

This proposal is using the technical approach which is the foundation for the SNL/NM corrective action process. The details of the SNL/NM technical approach are provided in Appendix C of the PIP. The first step in the technical approach is the data qualitative review step (the same step used to determine whether the SWMU is eligible for administrative NFA). Should significant uncertainties remain, the assessment of the SWMU continues within the SNL/NM technical approach.

At this site, sufficient data were not available to compare to established action levels or to develop site-specific action levels. Background soil samples were collected and analyzed to

develop upper tolerance limits (UTLs) for metals. Site-specific data were collected to compare to existing soil action levels (proposed Subpart S action levels) (EPA July 1990) and background UTLs. If site-specific concentrations exceeded the proposed Subpart S action levels or UTLs, then risk was analyzed. The site-specific concentrations were compared to the derived risk assessment action levels. Concentrations less than these action levels, either proposed Subpart S action levels, background UTLs, or derived risk-based values, triggered this NFA proposal for Site 50.

#### 1.3 Local Setting

SNL/NM occupies 2,829 acres of land owned by the Department of Energy (DOE), with an additional 14,920 acres of land provided by land-use permits with Kirtland Air Force Base (KAFB), the United States Forest Service (USFS), the State of New Mexico, and the Isleta Indian Reservation. SNL/NM has been involved in nuclear weapons research, component development, assembly, testing, and other nuclear activities since 1945.

ER Site 50 (Figure 1) is located on land owned by DOE. The site is situated east of the Technical Area (TA) II fence in a slight depression on top of the escarpment northwest of Tijeras Arroyo.

Surficial deposits in the SNL/NM KAFB area lie within four geomorphic provinces, which in turn contain nine geomorphic subprovinces. Site 50 lies within the Tijeras Arroyo subprovince. The Tijeras Arroyo subprovince is characterized by broad, west-sloping alluvial surfaces and the 50-meter-deep Tijeras Arroyo. The Tijeras Arroyo subprovince contains deposits derived from many sources, including granitic and sedimentary rocks of the Sandia Mountains, sedimentary and metamorphic rocks of the Manzanita Mountains, and sediments of the Upper Santa Fe Group.

#### 2. History of the SWMU

#### 2.1 Sources of Supporting Information

In support of this request for a risk-based NFA decision for ER Site 50, a background study was conducted to collect available and relevant site information. Interviews were conducted with SNL/NM staff and contractors familiar with site operational history.

The following information sources were available for the use in the evaluation of ER Site 50:

- Confirmatory sampling program conducted in September 1994
- Risk analysis for four metals and five radionuclides
- One surface radiation survey
- One unexploded ordnance/high explosives (UXO/HE) survey
- Reconnaissance Soil Sampling (DOE January 1989)
- Interviews and personnel correspondence
- Historical aerial photographs spanning 40 years
- The Comprehensive Environmental Assessment and Response Program (CEARP) Phase I (DOE September 1987)

#### 2.2 Previous Audits, Inspections, and Findings

ER Site 50 was first listed as a potential release site based on the CEARP interviews in 1985 (DOE September 1987), which noted that it was a rocket-powered centrifuge used to conduct acceleration tests on various weapon components. It was also noted in the CEARP that personnel do not recall any breakup of components that might have contaminated the area. However, there could be residual rocket propellants. The Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) finding was uncertain for Federal Facility Site Discovery and Identification Findings (FFSDIF), Preliminary Assessment, and Site Inspection; therefore, no Hazard Ranking System or Modified Hazard Ranking System migration mode scores were calculated for the SWMU (DOE September 1987). Site 50 was not included in the Environmental Protection Agency (EPA) RCRA Facility Assessment (RFA) in 1987 (EPA April 1987).

#### 2.3 Historical Operations

The Old Centrifuge was an outdoor rocket-propelled centrifuge that was used in the early 1950s to test units under G forces. The concrete centrifuge pad has a diameter of 80 to 90 feet. The site has a 7-foot high wooden retaining wall on the north, west, and south sides. The east side is open. The centrifuge arm assembly, which has a 20-foot radius, is sitting outside the wall to the north and appears to be intact. Control wiring to the center axis of the centrifuge was suspended from a cable between two telephone poles on the north and south side of the pad. The control wiring went to a bunker located to the southwest over the escarpment. The bunker had an electrical transformer containing polychlorinated biphenyl (PCB)-bearing oil. The electrical transformer has been removed. No spills or leaks were reported.

The centrifuge was driven by two solid-fuel rocket motors. The combustion byproducts produced by these rocket motors were carbon dioxide, carbon monoxide, water, hydrochloric acid, aluminum oxide, and possibly barium oxide. No high explosives are known or suspected at the site. The rocket orientation would expel combustion byproducts toward the retaining wall and the opening to the east.

#### 3. Evaluation of Relevant Evidence

#### 3.1 Unit Characteristics

The centrifuge pad has a diameter of 80-90 feet and is made of concrete. The site has a partial structural containment which consists of a 7-foot high wooden retaining wall on the north, west, and south sides. The east side is open.

#### 3.2 Operating Practices

The retaining wall limited the area in which the rockets would expel combustion byproducts. The rocket propellant was consumed in the rocket motor case. Under normal operating conditions, no unburned propellant would be released.

#### 3.3 Presence or Absence of Visual Evidence

There is no discoloration of soils along the open eastern end of the centrifuge and the concrete pad is not stained.

#### 3.4 Results of Previous Sampling/Surveys

In 1987, a reconnaissance investigation was conducted at the Old Centrifuge Site. Samples were analyzed for uranium, trinitrotoluene (TNT), hue, saturation, lightness (HSL) inorganic constituents, toxicity characteristic leaching procedure (TCLP) constituents, and environmental protection (EP) toxicity constituents. Metals, including barium, were detected at concentrations well below regulatory action levels. Total uranium concentrations were typical of area background levels. TNT, pesticides, PCBs, herbicides, and semivolatile TCLP compounds were not detected. Additional information on this sampling program is in Appendix E.

In 1994, the site was visually surveyed for surface indications of UXO/HE. No UXO/HE were found (SNL/NM 1994a). Also in 1994, a surface radiation survey was conducted on the entire site using an Eberline ESP-2 portable scaler, with an Eberline SPA-8 (2 inch X 2 inch sodium iodide) detector. A 30-second integrated count was performed at each proposed sample location, while scanning the detector over an area approximately 2 feet in radius around the sample location. The alarm was set at 1.3 times the background count rate. No alarms occurred during the survey. No surface anomalies were detected (SNL/NM 1994b).

#### 3.5 Assessment of Gaps in Information

Due to the potential constituents of concern (metals, specifically lead, beryllium, and barium; depleted uranium<sup>1</sup>; and high explosives), contamination, if any, is not expected to be at shallow depths. Therefore, a surface (0-6 inches) and shallow subsurface (6-36 inches) soil sampling program was developed and implemented in September 1994.

#### 3.6 Confirmatory Sampling

The Confirmatory Sampling and Analysis Plan (SAP) can be found in Appendix A. Those soil sample results exceeding an action level are summarized in Table 1. A complete list of "hits" or detections and quality assurance (QA) results can be found in Appendix B. For health and safety purposes, a photoionization detector, OVM, was used throughout the field program. The OVM measured no anomalous vapor concentrations.

<sup>&</sup>lt;sup>1</sup> These constituents are related to combustion products.

Four surface samples and four subsurface samples were collected at the most likely area for contamination, along the open eastern end (Figure 1). Every sample was analyzed for metals<sup>2</sup>. Six samples were analyzed for explosive residues. As a general check for radioactive constituents, three samples were analyzed for tritium and isotopic uranium. One sample was analyzed for isotopic plutonium and two samples were screened with in-house gamma spectroscopy.

#### 3.6.1 Background Samples for Metals and Radioactive Constituents

UTLs for background metals were calculated from analyses of 24 samples collected in the vicinity of the 11 sites discussed in the SAP (Appendix A). UTLs or background 95<sup>th</sup> percentiles for background radionuclides were calculated from samples collected throughout KAFB (IT 1994). A discussion of background calculations and supporting data and analyses are included in Appendices C and D.

#### 3.6.2 Explosive Residues

No explosive residues were detected.

#### 3.6.3 Metals

Mercury, selenium, and silver were not detected at Site 50. All other site concentrations for metals, except one analysis for lead, were below UTLs. The maximum local background value for beryllium was 0.53 milligrams per kilogram (mg/kg). Beryllium was not detected above 0.53 mg/kg at Site 50. For lead, Sample 50-02-A had a concentration of 25 mg/kg, compared to a UTL of 23.1 mg/kg. A RCRA Subpart S action level was not proposed for lead. However, a memorandum from an assistant administrator to EPA regional division directors does supply a risk-based action level for lead in soils of 400 parts per million (ppm) (mg/kg) (EPA 1994). The metals data indicate acceptable environmental or human health risk.

#### 3.6.4 Radionuclides

Plutonium-239/240 and plutonium-238 were not detected above the minimum detectable activity (MDA) at Site 50. Uranium-238 and uranium-234 were detected in three samples with activities ranging from 0.52 to 0.58 picocuries per gram (pCi/g) for uranium-238 and 0.53 to 0.66 pCi/g for uranium-234. Uranium-235/236 was detected in two samples at activities of 0.023 and 0.028 pCi/g. All activities for uranium-238, uranium-234, and uranium-235/236 are below the base-wide background 95th percentiles and the maximum activities of the six local background analyses. Thallium was not detected at Site 50.

<sup>&</sup>lt;sup>2</sup> Although the TAL metal analytes include calcium, magnesium, potassium, and sodium, these non-toxic, major cations are not included in the evaluation. They do not pose a significant environmental or human health risk regardless of concentration.

Cesium-137 was detected with in-house gamma spectroscopy screens for Samples 50-01-A and 50-03-A at activities of 1.66 pCi/g and 2.82 pCi/g, respectively, compared with a basewide background UTL of 0.92 pCi/g. Additional off-site analyses for cesium-137 on Samples 50-01-A and 50-03-A indicated lower activities. Tritium was detected in three samples at activities ranging from 0.020 to 0.038 pCi/g.

#### 3.6.5 Quality Assurance Results

As discussed in the Confirmatory Sampling and Analysis Plan (Appendix A), quality assurance samples, including field duplicates, trip blanks and rinsates, were collected as part of the 11-site sampling program. Analyses indicate that the field soil duplicates were comparable to the original soil sample results. The trip blanks and rinsates indicated no significant sampling contamination. QA results can be found in Appendix B. Level I and Level II data verification was conducted on all data, as described in the PIP (SNL/NM 1994).

#### 3.7 Risk Analysis

To further evaluate the site data for radionuclides with activities above the background UTL (cesium-137) or radionuclides without background UTLs (tritium), a risk assessment was performed for the combination of tritium and cesium-137, assuming the maximum detected activities.

The risk calculations were designed to produce conservatively large estimates of radioactive dose to counter uncertainties in the soil data. This approach facilitates the following decision regarding future activities at Site 50:

- If the conservative estimates based on the soil data result in an unacceptable dose (greater than 10 mrem/year), further investigation and/or remediation will be needed; or
- If the dose estimates are acceptable, the potential for health hazards at the site is extremely low, and further actions will not be needed.

Radionuclide doses were computed using methods and equations promulgated in proposed RCRA Subpart S documentation (EPA 1990). Accordingly, all calculations were based on the assumption that receptor doses from radionuclides result from ingestion of contaminated soil.

Calculation of radionuclide doses required values of dose conversion factors, which are used to convert radionuclide intakes (in units of pCi/year) into effective dose equivalents (in units of mrem/year). Published values of dose conversion factors (Gilbert et al., 1989) exist for tritium and cesium-137.

To assure that the computed doses were conservatively large, only the maximum observed activity of each constituent at a site was employed. To consider combined effects, a radiological dose was calculated as the sum of the individual doses.

Following proposed Subpart S methodology, the equation and parameter values used to calculate the summed radioactive dose were:

DOSE = 
$$\sum_{i}$$
 [DSR(i) x S(i)]

**(1)** 

where:

DOSE = total effective dose equivalent (mrem/yr);

DSR(I) = dose-to-soil concentration ratio for the i<sup>th</sup> radionuclide (mrem/yr)/(pCi/g), = I X DCF(I);

S(I) = soil concentration of the i<sup>th</sup> radionuclide (pCi/g);

I = soil ingestion rate = 0.2 g/day = 73 g/yr; and

DCF(I) = dose conversion factor for the i<sup>th</sup> radionuclide (mrem/pCi).

The PIP stipulates that, for the purpose of computing media action levels, the total radioactive dose at a site should not be greater than 10 mrem/year (SNL/NM 1994), which corresponds to a cancer risk of less that 10<sup>-6</sup> excess deaths.

The input and results of the risk calculations are presented in Table 2. The summed radioactive dose is less than 10 mrem/year. Therefore, the site is considered to be risk-free in terms of radionuclide contamination.

#### 3.8 Rationale for Pursuing a Risk-based NFA Decision

In September 1994, surface soil and shallow subsurface soil samples were collected at the open east side of the centrifuge, the most likely location for combustion byproducts from the rockets to settle and the most likely area to find contamination. SNL/NM is proposing a risk-based NFA because representative soil samples from ER Site 50 have concentrations less than action levels; either proposed Subpart S action levels, background UTLs, background 95<sup>th</sup> percentiles, or derived risk-based values.

#### In addition

- A site visit in 1993 and several visits in 1994 (August and September) by ER
  personnel confirmed no discoloration in the soils on the open east side of the
  centrifuge.
- In June 1994, a UXO/HE visual survey was conducted by KAFB Explosive Ordnance Division (EOD) and found no UXO/HE ordnance debris at Site 50 (SNL/NM 1994a).
- In September, 1994, as part of the surface soil sampling effort at Site 50, a surface radiation survey was conducted (SNL/NM 1994b). No surface anomalies were detected at Site 50.

#### 4. Conclusion

Based upon the evidence cited above, ER Site 50 has no releases of hazardous waste or hazardous constituents that pose a threat to human health and/or the environment. Therefore, ER Site 50 is recommended for an NFA determination.

#### 5. References

#### 5.1 ER Site References

Gilbert, T.L., C. Yu, Y.C. Yuan, A.J. Zielen, M.J. Jusko, and A. Wallo, 1989. Implementing Residual Radioactive Material Guidelines, A Supplement to U.S. Department of Energy Guidelines for Residual Radioactive Material at Formerly Utilized Sites Remedial Action Program and Surplus Facilities Management Program Sites, prepared by Argonne National Laboratory for U.S. Department of Energy, ANL/ES-160, DOE/CH/8901, 203 pp.

Guttman, I., 1970. Statistical Tolerance Regions: Classical and Bayesian. Hafner Publishing, Darien, Connecticut.

Internation Technology Corporation (IT), 1994. Background Concentrations of Constituents of Concern to the Sandia National Laboratories/New Mexico Environmental Restoration Project Phase II: Interim Report.

- U.S. Environmental Protection Agency (EPA), July 1994. Guidance on Residential Lead-Base Paint, Lead-Contaminated Dust, and Lead-Contaminated Soil, memorandum from Lynn R. Goldman, M.D., USEPA Assistant Administrator to EPA Division Regional Directors.
- U.S. Environmental Protection Agency (EPA), July 1990. "Corrective Action for Solid Waste Management Units (SWMU) at Hazardous Waste Management Facilities, Proposed Rule," *Federal Register*, Vol. 55, Title 40, Parts 264, 265, 270, and 271.
- U.S. Environmental Protection Agency (EPA), 1989. "Statistical Analysis of Ground-Water Monitoring Data at RCRA Facilities, Addendum to Interim Final Guidance," U.S. Environmental Protection Agency, Office of Solid Waste, Waste Management Division, Washington DC.
- U.S. Environmental Protection Agency (EPA), 1989. "Risk Assessment Guidance for Superfund: Volume I, Human Health Evaluation Manual (Part A)." Office of Emergency and Remedial Response, Washington, DC. 20460.

Sandia National Laboratories/New Mexico (SNL/NM), 1994a. "Unexploded Ordnance/High Explosives (UXO/HE) Visual Survey of ER Sites Final Report, Albuquerque, New Mexico," Sandia National Laboratories, Albuquerque, New Mexico.

Sandia National Laboratories/New Mexico (SNL/NM), 1994b. "Summary of Radiological Survey Results by SNL Dept. 7714 for ER Sites 50, 227, 229, 230-234," Sandia National Laboratories, Albuquerque, New Mexico.

#### 5.2 Reference Documents

Department of Energy (DOE), Albuquerque Operations Office, Environment and Health Division, Environmental Program Project Group, January 1989, Reconnaissance Data Report, Sandia National Laboratories Albuquerque."

Department of Energy (DOE), Albuquerque Operations Office, Environmental Safety and Health Division, Environmental Program Branch, September 1987, draft "Comprehensive Environmental Assessment and Response Program (CEARP) Phase I: Installation Assessment, Sandia National Laboratories, Albuquerque, New Mexico."

Sandia National Laboratories/New Mexico (SNL/NM), August 1994. Environmental Restoration Project Information Sheet for Site 50, Old Centrifuge Site, Sandia National Laboratories, Albuquerque, New Mexico.

Sandia National Laboratories/New Mexico (SNL/NM), February 1994. Draft "Program Implementation Plan for Albuquerque Potential Release Sites," Albuquerque, New Mexico.

U.S. Environmental Protection Agency (EPA), April 1987. "Final RCRA Facility Assessment Report of Solid Waste Management Units at Sandia National Laboratories, Albuquerque, New Mexico," Contract No. 68-01-7038, EPA Region VI.

U.S. Environmental Protection Agency (EPA), August 1993. "Module IV of RCRA Permit No. NM 5890110518, EPA Region VI," issued to Sandia National Laboratories, Albuquerque, New Mexico.

U.S. Environmental Protection Agency (EPA), August 1992. "Hazardous Waste Management Facility Permit No. NM5890110518, EPA Region VI," issued to Sandia National Laboratories, Albuquerque, New Mexico.

#### 5.3 Aerial Photographs

Ebert & Associates, Inc., November 1994, "Photo-Interpretation and Digital Mapping of ER Sites 7,16,45,228 from Sequential Historical Aerial Photographs."

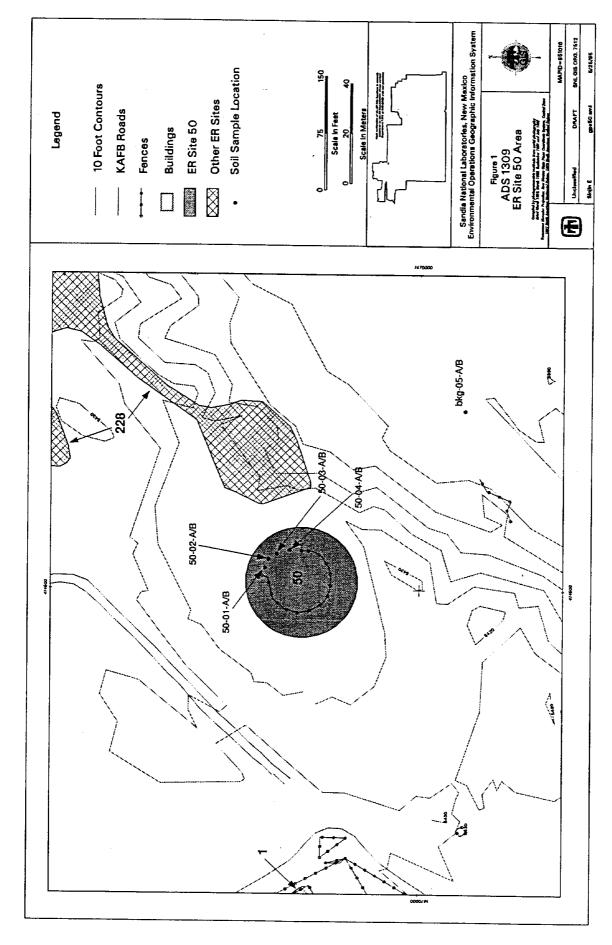


Figure 1. Old Centrifuge Site 50

Table 1. Site 50 Results of Shallow Soil Sampling and Analysis

Sample Identifier	Analytical Method	Constituent	Concentration <sup>1</sup>	Background <sup>2,3</sup>	Action Level <sup>4,5</sup>
50-02-A	TAL Metals (6010)	Lead	25	23.1	400
50-01-A	Tritium (600 906.0)	Tritium	0.024		36.9
50-01-B	Tritium (600 906.0)	Tritium	0.038		36.9
50-04-A	Tritium (600 906.0)	Tritium	0.02		36.9
50-01-A	Gamma Spec (in-house)	Cesium-137	1.66	0.92	274
50-01-A	Gamma Spec (in-house)	Cesium-137	2.82	0.92	274

#### <u>Notes</u>

Table 2. Risk Calculations for Site 50

Constituent	Activity (pCi/g)	DCF(I) (mrem/pCi)	Individual Dose (mrem/year)	Source of DCF
Cesium-137	2.82E+00	5.00E-05	1.03E-02	Gilbert et al., 1989
Tritium	3.80E-02	6.30E-08	1.75E-07	Gilbert et al., 1989
Summed Dose	•	·	1.03E-02	

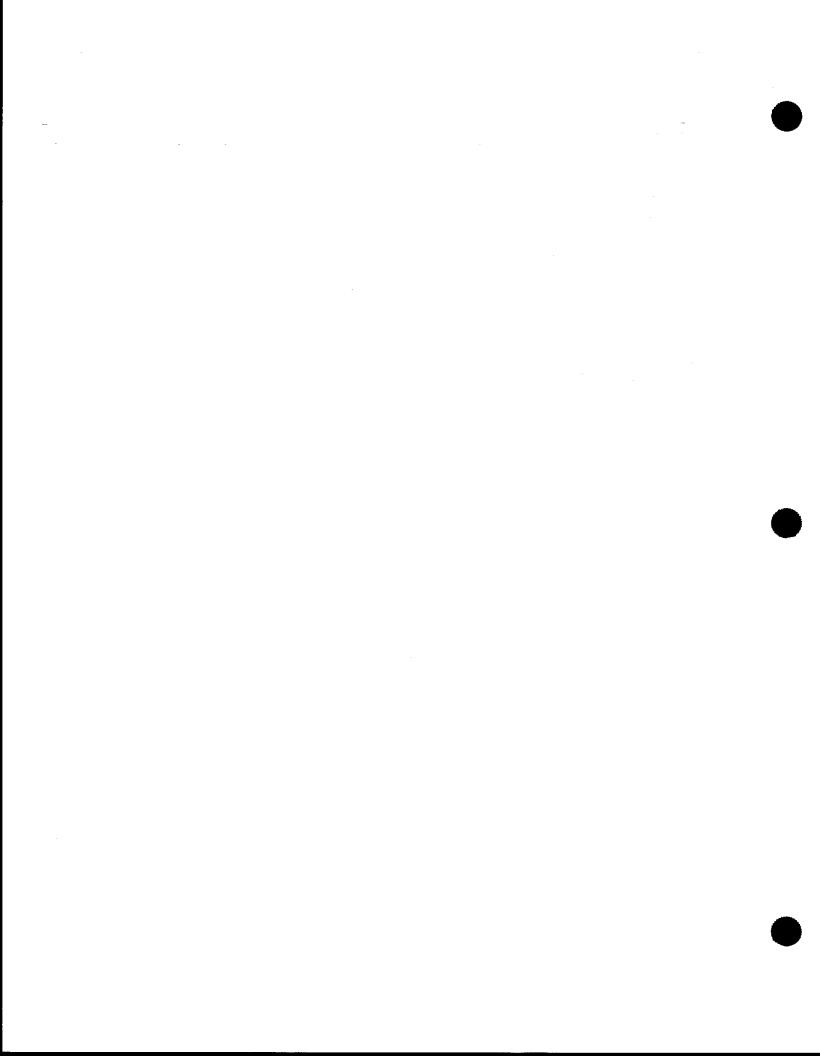
Lead concentrations are in mg/kg; tritium and cesium-137 activities are in pCi/g.

<sup>&</sup>lt;sup>2</sup>For lead, background is the 95 percent UTL for the local background data.

<sup>&</sup>lt;sup>3</sup>For cesium-137, background is the 95 percent UTL for the base-wide background data.

<sup>&</sup>lt;sup>4</sup>The lead action level is from EPA, 1994.

<sup>&</sup>lt;sup>5</sup>The tritium and cesium-137 action levels are calculated risk-based levels.



#### APPENDIX A

Confirmatory Sampling and Analysis Plan

#### APPENDIX B

**Analytical Results** 

#### APPENDIX C

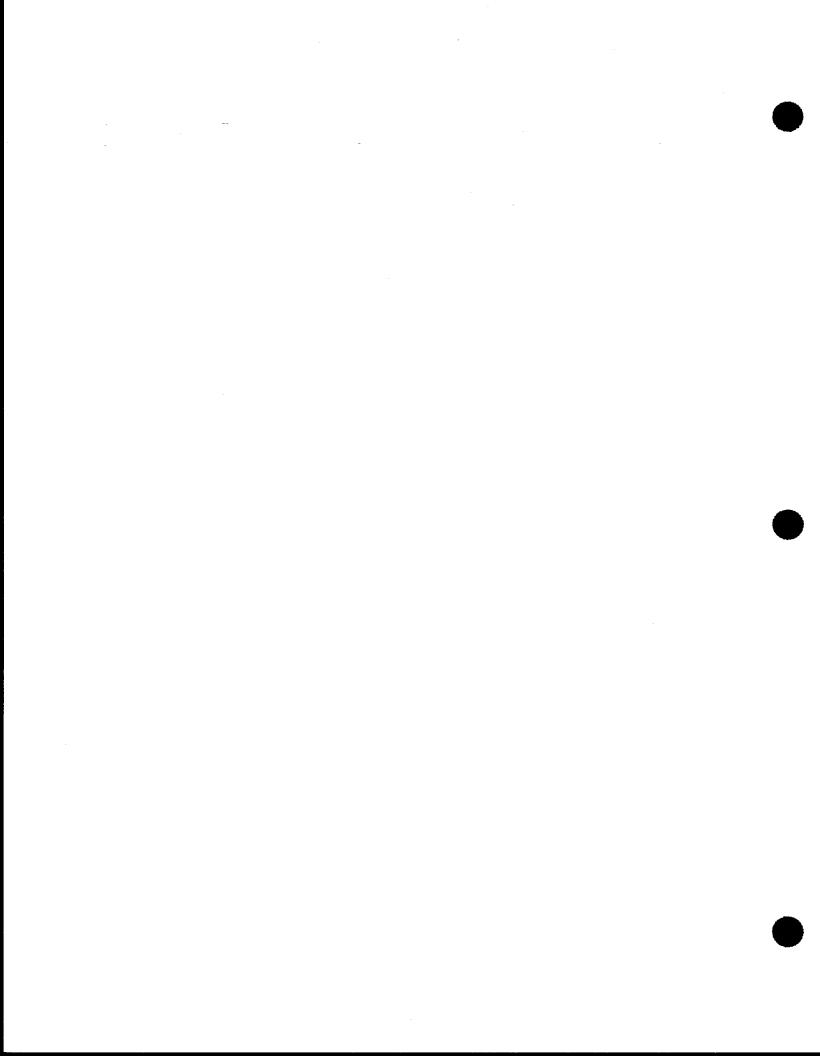
Background Calculations for Metals and Radionuclides

#### APPENDIX D

Probability Plots, Local Background UTL Calculations, and Base-wide Background UTLs for Radionuclides

#### APPENDIX E

Summary of Reconnaissance Data Report



#### Introduction

The purpose of the sampling and analysis described in this plan is to determine the appropriate way to proceed toward closure of 11 ( of the 17) sites in the Tijeras Arroyo Operable Unit. Based on the surface and shallow subsurface soil samples and analyses for the constituents of concern (COCs), one of three approaches will be pursued for each site:

- 1. A petition for "No Further Action" (NFA) will be produced for regulatory consideration;
- 2. A voluntary corrective measure (VCM) will be designed and implemented, hopefully followed by an NFA petition; or
- 3. The site assessment and eventual closure will follow the standard RFI/CMS path

Most of the sites covered by this Sampling and Analysis Plan (SAP) are outfalls from the storm water and sanitary sewer systems emanating from Sandia Technical Areas (TAs) I, II, and IV. The general sampling program for the outfalls will be to collect four samples at the head of the outfall, two samples of surface soil (0 to 6 inches deep) and two samples of shallow subsurface soil (18 to 36 inches deep) and four samples (two surface soil and two shallow subsurface soil) at the furthest extent of channel erosion and scour. The analytes for most of the samples are volatile organic compounds, semi-volatile organic compounds (BNAs), metals, chromium<sup>+6</sup>, for samples where chromium is found in a metals analysis, total petroleum hydrocarbon (TPH), explosives, Total Kjeldahl Nitrogen (TKN), nitrate/nitrite, and Gamma Spectroscopy for radionuclides, isotopic uranium, isotopic plutonium, tritium, and chlorodiphenyls (PCBs).

#### Sampling Procedures and Volumes

Surface soil samples will be collected with a stainless steel scoopula or trowel and placed in a stainless steel bowl. After at least 1000 ml<sup>1</sup> of soil has been collected, the soil will be thoroughly mixed in the bowl and transferred to two or three 500-ml sample bottles with a stainless steel scoopula. Sample bottles will be labeled accordingly and the appropriate sample information (sample depth, collection date and time, etc.) will be documented on the chain-of custody (COC) after each sample is collected. Samples will then be packaged and cooled to 4 degrees Celsius.

Shallow subsurface soil samples (18-36 inches) will be collected with a 2-inch (minimum) hand auger. A soil sample is collected by turning the auger clockwise and advancing it into the ground until the bucket at the end of the auger (last 6-8 inches) is full of soil or refusal occurs. Several runs with the auger is anticipated in order to obtain the appropriate volume. A hand shovel may also be used to bypass large rocks in order to continue with the auger. The auger is then extruded counter-clockwise from the ground and the soil is removed from the auger and placed in a stainless steel bowl. After 1,125² ml of soil has been collected, the soil will be mixed in the bowl and transferred to two or three 500-ml sample bottles and one 125-ml sample bottle with a stainless steel scoopula. Sample bottles will be labeled accordingly and the appropriate sample information will be documented on the COC after each sample is collected. Samples will then be packaged and cooled to 4 degrees Celsius.

#### Waste Generation and Equipment Decontamination

Decontamination of sampling equipment will be done between each sample.

Decontamination will include thoroughly washing the inside and outside of the sampling equipment with a spray of ALCONOX™ or LIQUINOX™ and water; rinsing with distilled,

<sup>&</sup>lt;sup>1</sup>The sample volume varies between 1,000 and 1,500 ml depending on the analyses for the sample.

<sup>&</sup>lt;sup>2</sup>The sample volume varies between 1,125 and 1,625 ml depending on the analyses for the sample.

deionized water; and drying before reusing. No soil waste will be generated. The soil removed from the hand-auger holes, while collecting samples at a depth of 18 to 36 inches, will be return to the hole. The sampling tools, which are scoopulas/trowels, hand-augers, and shovels, will be decontaminated with water and ALCONOX™ after each use. The decon leachate will be stored in capped 1-gallon containers. One or two containers will be used for each site and two to four containers will be used for the background samples. The containers will be labeled as "IDW" and the site number identified on each container. All the containers will be stored at Site 232, a central location. The leachate waste will be disposed according to the analytical results of the soil samples collected at the site.

#### Site Descriptions

The sites that will be sampled are

- Site 46, Old Acid Waste Line Outfall;
- Site 50, Old Centrifuge Site;
- Site 77, Oil Surface Impoundment;
- Site 227, Bldg. 904 outfall;
- Site 229, Storm Drain System Outfall;
- Site 230, Storm Drain System Outfall;
- Site 231, Storm Drain System Outfall;
- Site 232, Storm Drain System Outfall;
- Site 233, Storm Drain System Outfall;
- Site 234, Storm Drain System Outfall; and
- Site 235, Storm Drain System Outfall.

The site locations are shown in Figure 1. A description of the site history, conditions, previous investigations, and sampling plans are described in the following sections.

#### Site 46: Acid Waste Line Outfall

The Old Acid Waste Line carried wastes from several buildings in TA I. The waste line begins as a north-south trending, 750-feet long open trench in a grassy field northwest of Building 981-1 in TA IV. No pipe opening is visible at the "head" of the trench. As the trench crosses the field, it turns to the southeast and continues to a non-engineered spillway at the edge of Tijeras Arroyo. The spillway lies on a bank (40 to 50 feet of relief) composed of compacted alluvial sediment. Historical aerial photographs show vegetation, presumably supported by the discharge, growing southeast of the spillway to the active arroyo channel (about 200 feet distance from the spillway). The site is not restricted and is easily accessible.

During use, discharged effluent averaged an estimated 130,000 gallons per day. Use of the line has been discontinued. The line received wastes from plating, etching, and photo processing operations, and cooling tower "blow down". Acids and metals are target contaminants. Chromic acid and ferric chloride are mentioned specifically in the site history, and ferric chloride was found in the soils during a limited sampling event. Various radionuclides, possibly including tritium, uranium, and plutonium were used in TA I.

Building 863 was a source of discharge to the Acid Line. The information sheet for ER Site 98 (Building 863, TCA Photochemical Release: Silver Catch Boxes) indicates the presence of trichloromethane, silver, and photo-processing chemicals with an ammonia-like odor. The waste solution from the silver recovery unit reportedly was discharged to the Old Acid Waste Line, which is the only specific information about chemical discharges.

The site has been visually surveyed for surface indications of unexploded ordnance and high explosives (UXO/HE). No UXO/HE were found. Also, a surface radiation survey was

conducted on the entire site. No surface radiation anomalies were detected.

The sampling program includes four samples collected at the "head" of the site outfall (by the fire extinguisher training area west of TA IV) and four samples collected by the spillway into the Tijeras Arroyo drainage (Figure 1). Every sample will be analyzed for tritium, metals, chromium <sup>+6</sup> (if chromium is detected), TKN, and nitrate/nitrite. Half the samples will also be analyzed for semi-volatiles and cyanide. Additionally, all the subsurface samples will be analyzed for volatiles. The analytes are listed in Table 1. A "4" on the table indicates that ALL the samples will be analyzed

for that specific analyte whereas a "2" on the table indicates half the samples will have additional analyses for the analyte listed.

#### Site 50: Old Centrifuge

Site 50, Old Centrifuge, was an outdoor, rocket propelled centrifuge that was used in the early 1950s to test units under G forces. The facility is located east of the TA II fence in a slight depression on top the escarpment northwest of Tijeras Arroyo. The concrete centrifuge pad has a diameter of 80 to 90 feet. The site has a 7-foot high wooden retaining wall on the north, east, and south sides. The west side is open. The centrifuge arm assembly, which has a 20-foot radius, is sitting outside the wall to the north and appears to be intact. Control wiring to the center axis of the centrifuge was suspended from a cable between two telephone poles on the north and south side of the pad. The control wiring went to a bunker located to the southwest over the escarpment. The bunker had a electrical transformer containing PCB. The electrical transformer has been removed. The pad was not stained and no spills or leaks were reported.

The centrifuge was rocket driven by two T40 6-KS-3000 or two Deacon 3.5DS-5700 solid rocket motors. The combustion byproducts produced by these rocket motors were carbon dioxide, carbon monoxide, water, hydrochloric acid, aluminum oxide, and possibly barium oxide. No other HE is known or suspected at the site. The rocket orientation would expel combustion byproducts towards the retaining wall and the opening to the west. The rocket propellant would be consumed in the rocket motor case. Under normal operating conditions, no unburned propellant would be released.

In 1987, a reconnaissance investigation at five potential contaminated sites, including the Old Centrifuge Site, was conducted by the ER Project. Samples were analyzed for uranium, TNT, HSL inorganics, TCLP constituents, and EP Toxicity constituents. Metals, including barium, were detected at concentrations well below regulatory action levels. Total uranium concentrations were typical of area background levels. TNT, pesticides, PCBs, herbicides, and semi-volatiles TCLP compounds were not detected.

Prior to sampling, the surface will be surveyed for radiation. If contamination exists, it is expected to be around the edge of the centrifuge pad at the surface, probably along the open west side. The constituents of concern are metals (specifically lead, beryllium, and barium), depleted uranium, and high explosives. Four surface samples and four subsurface samples will be collected. The sampling locations will be biased toward the west side of the site because that is the open side (Figure 1). All surface samples will be analyzed for all the COCs. One-half of the subsurface samples will be analyzed for uranium and high explosives. All four subsurface samples will be analyzed for metals.

#### Site 77: Oil Surface Impoundment

The Oil Surface Impoundment Site is outside the TA IV fence, southeast of Building 981-1. The surface impoundment, which was constructed in the 1970's, is used to catch waste water from accelerators. At the time of the RCRA facilities environmental survey, the impoundment was unlined. Since then the impoundment was drained. Soil samples were analyzed for PCBs and

solvents. Based on the analytical results, the impoundment was determined to be clean. Subsequently, the impoundment was lined with geotextile and is now regulated under Sandia's Surface Water Discharge Program.

This site will not require UXO/HE or radiation surface surveys. Minimal confirmation sampling and analysis is proposed to verify that the site is clean. Three surface and three shallow subsurface samples are proposed. The samples will be collected along the perimeter of the existing lined pond (Figure 1). All the samples will be analyzed for PCBs. The subsurface soil samples also will be analyzed for volatile organic compounds (Table 1).

#### Site 227: Bunker 904 Outfall

Site 227 is an inactive outfall from the septic system for Building 904 (ER Site 48) in TA II. The site starts where the discharge exits the septic tank piping system, approximately 100 feet northeast of the southernmost point of TA II. The extent of the area influenced by the discharge may include the bank of Tijeras Arroyo below the outfall and some area between the outfall and the main channel of Tijeras Arroyo. The site is along the eastern edge of ER Site 45.

Building 904, built in 1948, was used for weapons assembly, HE testing, photo processing, and various other testing. Sanitary wastes were discharged to a septic tank, and other wastes were discharged to the outfall.

Mineral oil is also being considered a potential soil contaminant at all outfalls along the Tijeras Arroyo due to a recent release (June 1994) of mineral oil at Outfall 232 and vague historical records.

Possible soil contaminants are explosives, radioactive materials from weapons processing, including tritium, uranium, and plutonium, solvents (acetone, methylene chloride, methyl ethyl ketone, carbon tetrachloride, toluene, xylene, hexane, alcohols), and inorganics (ammonium hydroxide, barium, cadmium, silver, chromium, titanium, cyanide).

Access to this site is along the TA II perimeter road. This site is within the TA II testing exclusion zone. The best days to sample are generally Friday, Saturday, and Sunday, when testing ceases. Bruce Berry (telephone 845-8018) must be contacted to gain permission and access to this site. Prior to sampling

- tumbleweeds will be cleared from locations to be sampled and placed adjacent to the drainage;
- 2. these locations will be visually scanned for UXO/HE; and
- 3. these locations will be screened for surface radiation anomalies.

The proposed sampling program is to collect four surface soil samples and four shallow subsurface samples. Two surface and two subsurface samples will be collected at the outfall. The other two surface and two subsurface samples will be collected at the furthest visible channel erosion and scour (Figure 1). The analytes are listed in Table 1.

#### Sites 229 - 235: Storm Drain Systems Outfalls

These sites consist of the discharge areas at seven outfalls along the northern embankment of Tijeras Arroyo. The outfalls discharged industrial effluent and storm water from TAs I, II, and IV. Presently they only discharge storm water. The outfalls receive runoff from Site 96 (Storm Drain System) and other engineered drain systems within the three TAs. The sites are along approximately 3/2 miles of the embankment.

The specific constituents in the industrial effluent at these sites are not known. The possible discharged contaminants include chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, and other petroleum products. To cover this array of possible contaminants, soil samples will be analyzed for volatiles (subsurface samples only), semi-volatiles, metals and chromium of chromium is found in the metals analysis.

Mineral oil is also being considered a potential soil contaminant at all outfalls along the Tijeras Arroyo due to a recent release (June '94) of mineral oil at Outfall 232 and vague historical records. Therefore, soil samples will also be analyzed for TPH.

At Sites 229 through 234, prior to sampling

- 1. tumbleweeds will be cleared from locations to be sampled and placed adjacent to the drainage;
- 2. these locations will be visually scanned for UXO/HE; and
- 3. these locations will be screened for surface radiation anomalies.

Site 229 is due east of the footings of the old guard tower and the south "corner" of the TA II fence. It discharges near the top of the embankment through the center of ER Site 45. Access to this site is along the TA II perimeter road. This site is within the TA II testing exclusion zone. The best days to sample are generally Friday, Saturday, and Sunday, when testing ceases. Bruce Berry (telephone 845-8018) must be contacted to gain permission and access to this site. Because this site discharges from TA II, various radionuclides, possibly including tritium, uranium, and plutonium are of concern. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 230 is west of Building 970 in TA IV. A drain pipe discharges into a bowl-shaped concrete structure adjacent to Building 970A. Flow from this structure is directed to a drain and flume located approximately 120 feet further west. The flume carries the flow to a discharge point slightly above the base of the arroyo embankment. Doug Bloomquist (845-7455) must be contacted to ensure that no laser testing is being performed in the area. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 231 is west of Building 970 in TA IV. A drain pipe discharges to a concrete flume near the top of the embankment. The flume carries the flow to a discharge point near the base of the slope. Doug Bloomquist (845-7455) must be contacted to ensure that no laser testing is being performed in the area.. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 232 consists of two outfalls. One outfall is south of Building 970A, east of the lined lagoon. A drain pipe discharges to a concrete flume near the top of the embankment. The flume carries the flow to at discharge point near the bottom of hillside. On June 1, 1994, about 150 to 350 gallons of mineral oil was spilled into this outfall through the storm water drain by building 986. The day after the spill the site was screened for radiation and UXO/HE. No surface radiation anomalies or UXO/HE were found. Also, four surface soil and four subsurface soil samples were collected. The samples were sent to Quintera Laboratory in Denver for analysis for organics, metals, chromium<sup>16</sup>, and gamma spec. Other than TPH from the mineral, no contaminants were detected. A Voluntary Corrective Measure was conducted in July and August to remove soil contaminated with mineral oil above 100 mg/kg of TPH.

The second outfall in Site 232 also is south of Building 970A, west of lined lagoon, and approximately 120 feet east of the other Site 232 outfall. Discharge occurs from a concrete structure opening near base of embankment. Access to the site is along the road outside the south side of TA IV. Four surface soil and four subsurface soil samples will be collected at this drainage Figure 1). The analytes are listed in Table 1.

Site 233 is south-southwest of Building 986. Near the top of an escarpment, a small metal drain pipe discharges to an open drain which directs flow within another pipe before discharging near the base of the hillslope. Access to the site is along the road outside the south side of TA IV. Four surface soil and four subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 234 is southeast of Building 981I (Inflatable Building) and a lagoon impoundment (Site 77).

The site discharges into a steep-sided, deeply incised channel cut into the hillside. The drainage channel splits directly uphill of a tree. Access to the site is along the road outside the south side of TA IV. Both channels will be sampled. Six surface soil and six subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

Site 235 is immediately downstream of a large concrete spillway on the northeast side of Pennsylvania and south of the Skeet Range, at the point where the road comes off the north bank of the arroyo and descends into the channel. The flow moves in a confined channel after dropping down the spillway. The site has been cleared for visible surface UXO/HE and screened for surface radiation with no anomalies detected. This channel is considerably larger than the other outfall sites. Six surface soil and six subsurface soil samples will be collected at this site (Figure 1). The analytes are listed in Table 1.

#### **Background**

Background soil concentrations for organic contaminants should be negligible. Background concentrations for total metals and radionuclides must be determined for comparison to concentrations found at the sites. Twelve locations have been identified to collect samples for background determination (Figure 1). At each of these sites, one sample will be collected at a depth of 0-6 inches and a second sample collected at 18-36 inches (Table 1).. In addition, the background study report prepared by International Technology Corporation (May 1994) will also be used to evaluate the data.

#### **Quality Assurance**

As shown in Table 1, quality assurance samples will include the following:

- Field "duplicates" on more than 10 percent of the samples. These samples will be collected adjacent to the original surface soil sample and in the same hole as the original subsurface soil sample;
- Field soil blanks for more than 10 percent of the VOC analyses. These sample will be obtained from Sample Management Office (SMO) and will contain no VOCs; and
- One rinsate blank. All rinsate will be composited in one container. A sample of the
  rinsate will be analyzed for all constituents. The disposal method for the rinsate will be
  determined by the analytical results on this sample.

Soils	(0.90 906.0) mulino (0.00 7-79-081)	<del></del>				7	$\neg$				<del> </del>		4	"	+	Ш	7	1
Soils	(0.30e 003) multin				_1	, ,,	l l	- 1	ľ	- 1	Ī		- 10	าไ	- 1	ιı	9	1
Soils		T 4	_ -	-	4	4					<del>- </del> -			╁	+	Н	16	
Soils	CBs (8080)	4		1	r							+	+	1.	+-	Н	5 1	
Soils	1.109 000 (off-site) 500 901.1			T						<b>-</b>		╅──-	+	╁	╁	Н	-	ĺ
Soils	Samma Spec (In-House) 600 901.1	4	T	Ī	4	4	7	7	7	7	7	7	15	╬	╀	┦	9	
Soil	7.636) SON/cOV	4		T	4	1	+-	<b></b> -	+	+		+-``	+	╁	+	┦	36	
	LKN (sciq qiđeslion)		7	+	4	+	+-	+	<del> </del> -	-	<del> </del>		+	<u> </u>	1		6	
ខ្ល	Explosives Res (8330)		7	1	2	<del> </del>	+	+		+	<del></del>	┿	4	╀	_		6	ì
Sur	(3108) H9T		十	t	4	4	+ 4	4	+_	+			$\bot$	上	+		9	
Subsurface	Cyanide (acid digestion)		╁	╁	- 7	+	<del>  `</del>	<del>-</del>	4	4	Ψ	4	$\perp$		1		8	
	C <sub>1***</sub> (aqueous leaching)		╁	╁	4	4	<del> </del>		-	-			$\perp$	Ŀ			ല	
	(000\7\0\0) sisteM JAT		4	+	4		4	4	4	4	9	4	$\perp$	4	Ц		42	
	(000510103) sleteM_IAT	<u> </u>	╀	╁		4	4	4	4	4	9	4	12	ြ		-[	8	
	(07S8) sAN8	2			6	24	8	7	7	7	<sub>6</sub>	7		2		Ī	2	
	AOC2 (8540)	4		4	4	4	4	4	4	4	9	4		2	2	-	2	23
Ц	Number of Samples		4	4	4	4	4	4	4	+	9	4	12	⊢	H		┛	ı
	Isotopic Uranium (HASL-300 4.5)	2	2		7	8	-	+-	-	+-	+-	<del>                                     </del>	+	-	Н	_	3	ᆜ
	(180-67-7 000) muinotul9 ciqotosi	7	-		8	2	+-	+-	+-	<del> </del>	<del>                                     </del>	-	╌	7	Н		_	3
	(0.30e 003) muithT	4	2		4	4	~	- 7	8	-	<del>                                     </del>	<del> </del>	<u>اس</u>	₩.	Ц	_		26
Γ	PCBs (8080)		Н	4		<del>                                     </del>		<del>                                     </del>	+	2	7	7	<u>س</u>	Ц	$\sqcup$		_	46
Γ	Gamma Spec (Off-site) 600 901.1	7	H		2	~	<del> </del>		<del> </del>	-	<del> </del>	<u> </u>	╄		Ц	-	-	듸
ſ	Gamma Spec (In-House) 600 901.1	4	2	Н	4	4	2	7	<del> </del>	-			L	٦				∞]
Solls		4	Н	$\dashv$	4	-			"	7	7	2	12	Ц			_	?
ဖွဲ့	NO <sub>2</sub> /NO <sub>2</sub> (353.2)	4	H	$\dashv$	·····	ļ		<u> </u>	ļ	<u> </u>				-	_	- (4)	2   9	2
8	TKN (scid digestion)			$\dashv$	4									$\vdash$	1	- 5	? (	2
	Explosives Res (8330)		4	$\dashv$	- 7	ļ	<u> </u>		<u> </u>	<u></u> .				F	7	- 0	, ;	2
7	TPH (8015)		Ц	4		4	4	4	4	4	9	4		4	1	- 12	:	ब
-	Cyanide (acid digestion)	- 7	Ц	4	. (1)									-	٦,	- 4	1	=
F	Cr*** (aqueous leaching)	4	4	4	4	4	4	4	4	4	9	4		4	٦,	- 5	_	1
L	(0007/0103) slataM JAT	4	4		4	4	4	4	4	4	မ	4	12	2	1	- 2	-4-	7
Ļ	(OTSB) sANB	2	T	$\top$	7	7	2	7	2	- 2	9	2	H	7	+	3	1 27	<u>. 1</u> :
	Number of Samples	4	4.	7	4	4							$\dashv$	-	+	_	4.	$\exists$
$\dagger$			+	4			4	4	4	4	9		12		1	8	116	
	Potential Contaminants Ferric chloride chronic and other acids		Convert propellant and residues	-	High explosives, radioactive materials, nitrate, toluene, methanol, other solvents, carbon tetrachloride, ammonium hydroxide, barlum, cadmium, silver, chromium, titanium, cyanide	Chromates, antifoulants, chromium, sadium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromlum, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products	Chromates, antifoulants, chromium, sodium hydroxide, hydrochloric acid, chromosulfuric acid, diesel, other petroleum products		Sa C	as Z	Totals	Totals - Surface Plus Subsurface	* Analyze for Cr*6 only if Cr is defended in motel
	Site Name	Outfall (Tijeras Arroyo)	Oil Surface Impoundment	in the state of th	Bidg. 904 outfall (TA-2)	Storm Drain System Outfall	Background	Field Soil Blank	Rinsate		Totals - 5							
	Site	46	-	+	227	229	230	231	232	233	234	235	e S	Y O	i S	_	_	1

-		

# Appendix B Analytical Results

-	-		

			,					, _										
	Nickel	5.2	4.5	7.1	4	4.4	4.1	2.7	3	46S-muins1U	0.7	0.0					0.5	
	Мегсигу	Q.	S	Q	QN N	QN	SN N	QN	ND	852\2557muine1U	0.03	0.023					0.028	
	Manganese	230	130	210	150	170	140	140	120	862-muina1U	9.0	0.0					0.5	1-
	muisəngeM	3800	3000	4400	2100	2600	2600	3000	2500	86S-muinotul9	00'0							
	reaq	22	6.6	25	7.5	8.4	2.5	4.8	5.4	04S\68S-muinotul9	0.00							
esults	lron	6100	7600	6700	7100	7800	2500	6200	4900	muithT	0.02	0.04			<b></b> -		0.02	
Soil Results	Copper	11	6.2	12	2.3	8.1	6.6	4.6	4.4	781-muisəO	1.8						Q.	П
Site 50	tladoO	4.4	3.9	4.3	3.7	3.8	3.7	3.3	2.9	∑£1-muisə⊃	1.7				2.8			
S	Сһготіит	4	4	വ	က	4	4	3	2	Explosives Res.	Q.	ND	<del>Q</del>		QN	2	g	П
	muiolsO	96000	46000	120000	21000	28000	47000	49000	41000	Sinc	42	18	69	20	24	9.9	16	14
	muimbeO	9.1	- 3	9.		1.3	1.2	1.1	0.7	muibensV	15	17	11	13	15	16	19	13
	Beryllium	0.4	0.3	0.4	2	0.3	呈	S	2	muillsdT	S	2	ND	QN	S	QN	DN	ND
	muisa	210	110	220	59	82	110	96	91	muibo2	2	S	QN	Q.	S	Q.	QN	QN
	oinəsıA	ω	7	4	7	4	2	5	က	Silver	일	9	QN N	<u>S</u>	일	S	Q.	QN
	YnomitnA	6	_	12	ည		8	6	2	muinələ2	밁	위	2	-	9	S	9 P	QN
	munimulA	4000	3100	5800	2900	3600	3200	2600	2100	muissetoq	1300	7	1300	ᅱ	1		-	510
	Sample Identifier	50-01-A	50-01-B	50-02-A	50-02-B	50-03-A	50-03-B	50-04-A	50-04-B		50-01-A	50-01-B	50-02-A	50-02-B	50-03-A	50-03-B	50-04-A	50-04-B

Concentrations in mg/kg
Activities in pCi/g
Sample Identifier XX-XX-A - surface soil samples
Sample Identifier XX-XX-B - subsurface soil samples

p2 ....

	HdT							9	84						Γ								QN QN
	total-Xylenes							-															0.001
	Styrene																					0.001 J	
	Ругеле	0.040						0.19 J	0.28 J														
	РћепалИтепе	0.055 J	0.051 J					0.17 J	0.18 J														
	Methylene Chloride																			0.003			
lituents	Fluoranthene	0.066 J	0.038 J					0.23 J	0.20 J														
c Const	Di-n-octyl phthalate													0.16 J									
Organi	Сһгуѕепе							0.11 J	0.12 J														
ilts for	Benzo(b)fluoranthene							0.16 J	0.16 J														
ice Resu	Benzo(a)pyrene							0.050 J	0.092 J														
ality Assurance Results for Organic Constituents	Benzo(a)anthracene							0.071 J	0.006														
Quality	ənotəəA				0.006 J													0.019	0.015			0.015	0.010
	4-Methyl-2-pentanone			0.001 J														0.002 J					
	S-Hexanone																	0.003 J					
	S-Butanone			ቦ 200'0	r 900'0	0.004 J	0.005 J			0.006 J	0.006 J	0.006 J	0.006 J	0.003 JB		0.006 JB	0.004 JB	0.010 B	0.009 JB	0.004 JB	0.007 JB	0.007 JB	0.005 JB
	Sample Type	original	duplicate	trip blank	trip blank	trip blank		trip blank 0.007 JB	rinsate														
	Sample Identifier	227-01-A	227-01-A	227-01-B	227-01-B	227-04-B	227-04-B	229-01-A	229-01-A	229-02-B	229-02-B	229-03-B	229-03-B	230-04-B	230-04-B	235-02-B	_	Site 227	Site 229	_	Site 232		Site 235

'	Quality Assurance Results for Inorganic and Radiological Constituents																
Sample Identifier	Sample Type	Aluminum	Antimony	Arsenic	Barium	Beryllium	Cadmium	Chromium	Cobalt	Copper	Iron	Lead	Manganese	Mercury	Nickel	Vanadium	Zinc
227-02-A		5800	9.3	5.9	180	ND	2.1	6.6	4.1	7.8	13000	7.5	160	ND	5.4	27	51
227-02-A	duplicate		11	1.4	150	0.25	2.5	6.4	4.1	13	14000	9.1	170	ND	5.9	28	51
227-03-B		5100		0.92	140	ND	2.1	5.9	4.5	11	13000	7.5	200	ND	5.4	25	48
227-03-B			9.9	5.6	140	0.25	2.9	7.4	4.6	10	16000	8.9	230	ND	5.9	33	50
229-04-A		8100	13	5.7	150	0.32	2.3	8.0	4.2	7.9	13000	12	210	ND	6.3	24	55
	duplicate			1.5	140	0.30	2.2	8.0	4.2	7.7	12000	11	190	ND	6.2	24	52
230-04-B		1500	3.3	1.6	130	ND	0.61	2.3	ND	18	3500	4.2	110	ND	3.0	9.1	82
230-04-B			4.9	1.7	140	ND	0.68	3.1	2.5	15	4500	4.1	120	ND	3.4	9.7	71
235-01-A		3600	6.2	5.1	150	ND	2.7	6.0	8.4	6.6	20000	7.6	210	ND	4.5	36	66
235-01-A	duplicate		5.3	1.3	160	ND	1.6	4.2	5.7	6.5	12000	9.4	180	ND	4.4	22	66
50-01-B	original	3100	6.5	2.1	110	0.25	1.3	4.1	3.9	6.2	7600	6.6	130	ND	4.5	17	18
50-01-B	duplicate		7.5	2.0	110	0.26	1.3	4.3	4.0	5.7	8800	5.9	150	ND	4.2	18	21
50-02-A	original	5800	12	4.2	220	0.38	1.6	5.2	4.3	12	6700	25	210	ND	7.1	11	69
50-02-A	duplicate	7000	14	6.4	280	0.55	2.2	8.3	6.1	17	9000	35	290	0.04	9.4	18	61
Bkg-05-A	original	6400	13	5.7	210	0.53	1.8	6.1	6.6	14	10000	16	330	ND	8.9	22	37

7.6

ND

Bkg-05-A duplicate 5900 12

rinsate

ND

ND

Site 235

ND

190 0.50

ND

1.7

ND

6.0

ND

6.3

ND

14

ND

											Notes on Quality Assurance Data
Sample Identifier	Sample Type	TKN	NO <sub>3</sub> /NO <sub>2</sub>	Potassium 40	Lead 212	Lead 214	Plutonium 239/240	Uranium 238	Uranium 235/236	Uranium 234	Explosive residues were not detected in Site 50 duplicate sample  Hexavalent chromium was not detected in five duplicates and one decon rinsate
227-02-A		400	2.7								Cyanide was not detected in two
227-02-A		320	9.3						_		duplicates and one decon rinsate
227-03-A		_					0.004	0.4	0.15	0.61	888
227-03-A								0.67	0.023	0.67	PCBs were not detected in one Site 77
227-03-B								0.72	0.11	0.72	duplicate sample
227-03-B		220	ND				-				Tritium and Plutonium-238 were not
227-03-B				27.8	0.71	0.7					detected in four duplicate samples
227-03-B		190	1.4								and a spinoate samples
229-01-A							0.007	0:45	0.17	0.67	Selenium, silver, and thallium were not
229-01-A	duplicate							0.73	0.034	0.6	detected in any quality assurance
229-03-B								0.45	0.058	0.45	samples
229-03 <b>-</b> B	duplicate							0.99	0.06	1	
				_							

ND ND ND ND ND ND

24 | 36

10000 16 320 ND 8.7

# Appendix C Background Calculations for Metals and Radionuclides

#### Appendix C. Background Calculations for Metals and Radionuclides

To evaluate metals data, 24 background samples were collected for metals analyses.<sup>4</sup> Distribution analyses was performed first by constructing histograms. The histograms indicated a parametric distribution. Outliers were screened in a two-step process as described in the base wide background report (IT 1994). The first step is to perform an "a priori" screening for very high values relative to the rest of the data set. This is qualitatively performed by visually examining a column of sorted values. Maximum values that are a factor of 3 or 4 times higher than their nearest neighbor are removed from the data set during this step. None of the anomalous values were deleted by the "a priori" process.

The second step, from EPA, 1989, determines whether an observation that appears extreme fits the data distribution. A statistical parameter,  $T_n$  is calculated:

$$T_0 = (X_0 - X_a)/S$$

where:

 $X_n$  = questionable observation;

X<sub>a</sub> = sample arithmetic mean; and

S = sample standard deviation

 $T_n$  is compared to a table of one-sided critical values for the appropriate significance level (upper 5 percent) and sample size from a table provided in EPA 1989. Extreme concentrations for barium, calcium, chromium, copper and nickel were identified as outliers and were excluded from the data set. These anomalous values may have resulted from laboratory or sampling error.

Probability plots were then replotted to determine whether the data fit normal or lognormal populations. These plots are shown in Appendix D. The UTL<sup>5</sup> was calculated for data sets that fit a normal or lognormal distribution. Data sets are provided in Appendix D. As recommended by EPA, a tolerance coefficient value of 95 percent was used (EPA 1989). Most metals background data fit lognormal distributions. Iron and zinc data fit normal distributions. UTLs were not calculated for mercury, selenium, and silver because mercury and selenium were not detected and silver was detected only once in the 24 background samples. The beryllium background data did not fit a normal or lognormal distribution. The maximum value in a data set is commonly taken as the UTL in a non-parametric setting (Guttman, 1970). The maximum background beryllium concentration was 0.53 mg/kg.

Base-wide background UTLs for radionuclides were established by International Technology (IT) Corporation to compare and evaluate radionuclide data (IT, 1994). A table is provided in Appendix

<sup>&</sup>lt;sup>2</sup>These data are referred to as local background data. The data collected throughout Kirtland Air Force Base (KAFB), with most of the data collected within SNL/NM technical areas, are called base-wide background data (IT 1994).

 $<sup>^{3}</sup>$ UTL = x + K•S, where:

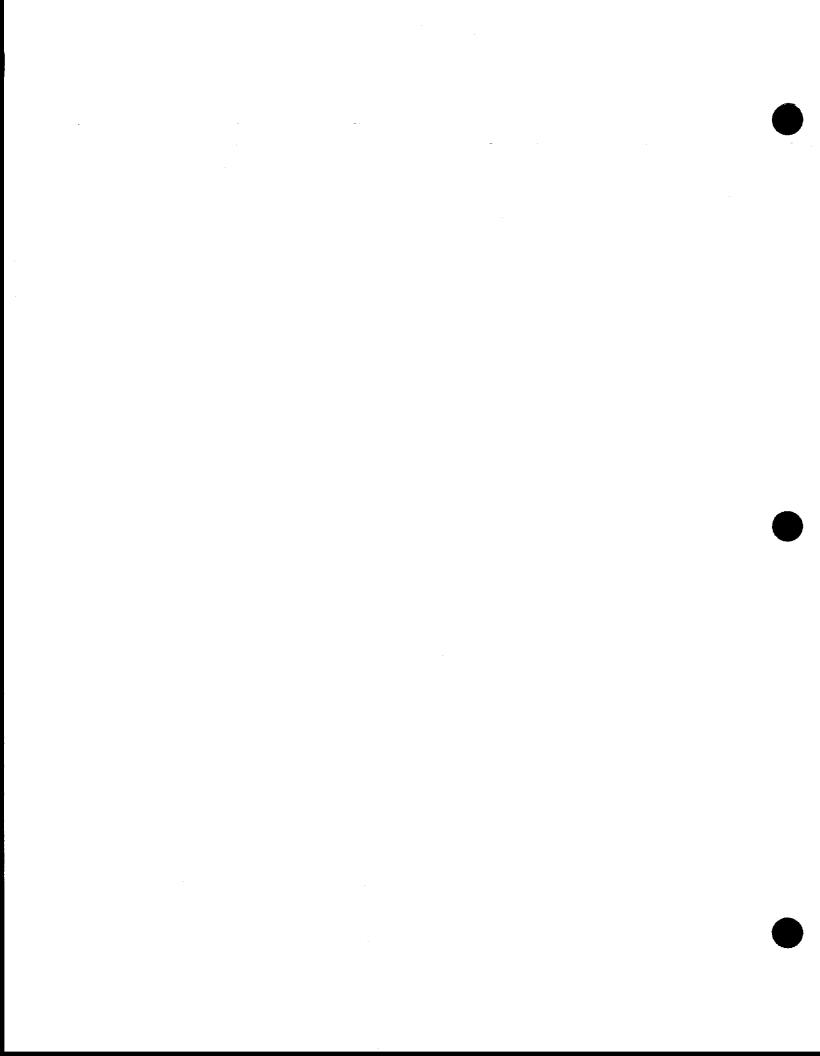
UTL = Upper tolerance limit;

x = Sample arithmetic mean (for normal distribution), sample geometric mean (for lognormal distribution);

S = Sample standard deviation; and

K = One-sided normal tolerance factor (95 percent for these evaluations).

D with radionuclide background data and the corresponding UTLs. The maximum activity from the six local background samples for isotopic plutonium and isotopic uranium was used as an additional method to evaluate the data. Also, in-house gamma spectroscopy was performed on all 24 background samples and indicated low levels of radioactivity but no significant contamination.

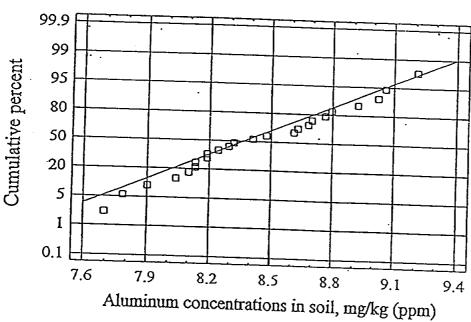


# Appendix D Probability Plots, Local Background UTL Calculations, and BaseWide Background UTLs for Radionuclides

Summary Statistics for log(Aluminum)

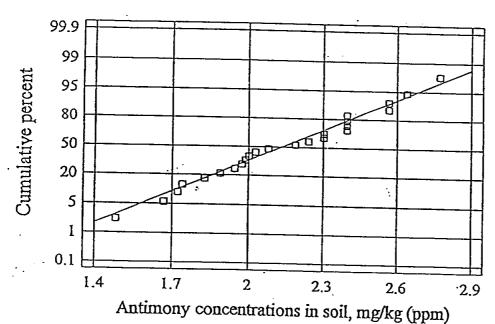
Count = 24 Average = 0.42942 Median = 8.36529 Yode = Jeometric mean = 8.41976 /ariance = 0.170246 Standard deviation = 0.412609 Standard error = 0.0842235 finimum = 7.69621 faximum - 9.21034 lange = 1.51413 ower quartile = 8.13153 pper quartile = 8.73178 interquartile range = 0.600253 kewness = 0.132255 itnd. skewness = 0.26451 urtosis = -0.792361 tnd. kurtosis = -0.792361 oeff. of variation = 4.89487 um = 202.306

# Lognormal Probability Plot for Aluminum



ry Statistics for log(Antimony) 'erage = 2.14609 dian = 2.13275 de = 2.3979 ometric mean = 2.12004 riance = 0.113831 andard deviation = 0.337389 andard orror = 0.0688692 nimum = 1.4816 ximum = 2.77259nge = 1.29098 wer quartile = 1.91649 per quartile = 2.3979 cerquartile range = 0.481405 wness = -0.040772 id. skewness = -0.0815441tosis = -0.744171 nd. kurtosis = -0.744171 of variation = 15.7211 i = 51.5062

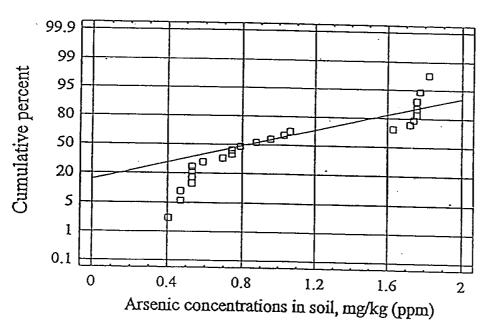
# Lognormal Probability Plot for Antimony



Summary Statistics for log(Arsenic)

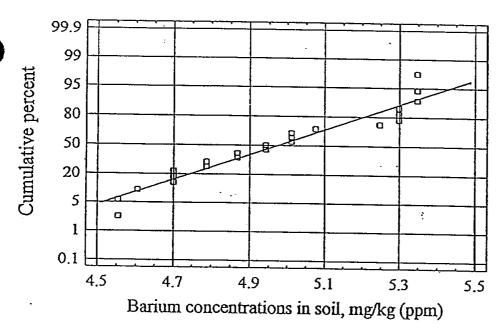
Count = 24 Average = 1.038 Median = 0.031963Mode = Geometric mean = 0.908119 Variance = 0.291153 Standard deviation = 0.539586 Standard error = 0.110143 Minimum - 0.405465 Max1mum = 1.82455 Range = 1.41908 Lower quartile = 0.530628 Upper quartile = 1.73162 Interquartile range = 1.20099 3kewness = 0.463036 Stnd. skewness = 0.926071 (urtosis = -1.58507 Stnd. kurtosis = -1.58507 Coeff. of variation = 51.983 jum = 24.9121

#### Lognormal Probability Plot for Arsenic



mary Statistics for log(Bacium) **4.96948** ian = 4.94164 · = 5.34711 netric mean = 4.96236 ance = 0.0740602 idard deviation = 0.27214 dard error = 0.0567451 mum = 4.55388 .mum = 5.34711 e = 0.793231 r quartile = 4.70048 r quartile = 5.29832 rquartile range = 0.597837 ness = 0.0653415 . skewness = 0.127931 osis = -1.30542. kurtosis = -1.27794 f. of variation = 5.47622**= 114.298** 

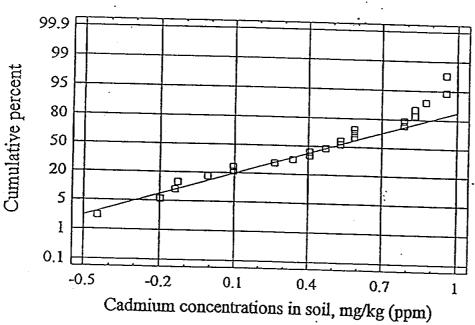
#### Lognormal Probability Plot for Barium



Summary Statistics for log(Cadmium)

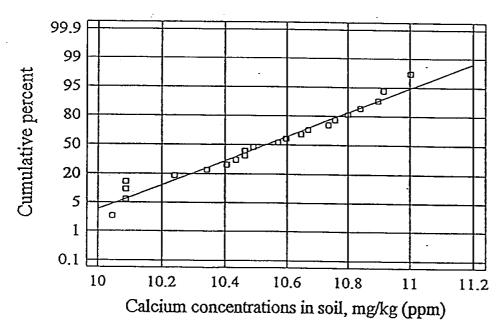
Count = 24 Average = 0.416764 Median = 0.500316 Mode = Geometric mean = Variance = 0.159937 Standard deviation = 0.399922 Standard error = 0.0816337 Minimum = -0.446287 Maximum = 0.955511 Range = 1.4018 Lower quartile = 0.0953102 Upper quartile = 0.788457 Interquartile range = 0.693147 Skewness = -0.506707Stnd. skewness = -1.01341Kurtosis = -0.674504 Stnd. kurtosis = -0.674504Doeff. of variation = 95.9587 3um = 10.0023

# Lognormal Probability Plot for Cadmium



mmary Statistics for log (Calcium) = 10.5579 dian = 10.5713 de = 10.0058ometric mean = 10.5532 riance = 0.10513 andard deviation = 0.324237 andard error = 0.0676081 nimum = 10.0432ximum = 11.2645 nge = 1.22121 wer quartile = 10.3417 per quartile = 10.7996 terquartile range = 0.457833 wness = 0.109797 nd. skewness = 0.214971tosis = -0.415646 id. kurtosis = -0.406895 of variation = 3.07103 x = 242.832

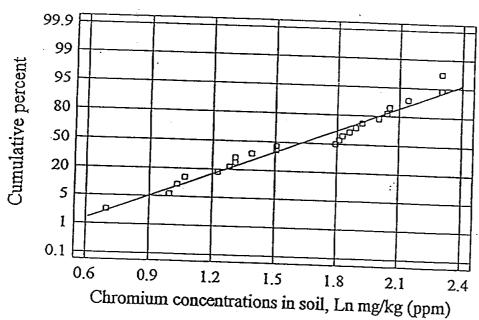
#### Lognormal Probability Plot for Calcium



Summary Statistics for log(Chromium)

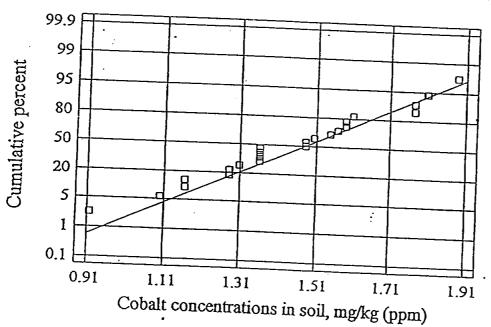
Count = 23 -Average = 1.61041 Median = 1.79176 Mode = Geometric mean ≈ 1.55042 Variance = 0.204195 Standard deviation = 0.451879 Standard error = 0.0942233 finimum = 0.693147 faximum = 2.30259 lange = 1.60944 ower quartile = 1.28093 oper quartile = 2.00148 nterquartile range = 0.720546 kewness = -0.274151 tnd. skewness = -0.536757urtosis = -0.905395 tnd. kuctosis = -0.886332 oeff. of variation = 27.9211 um = 37.2235

# Lognormal Probability Plot for Chromium



umunary Statistics for log(Cobalt) gc = 1.29969 edian = 1.42129 cometric mean = iriance = 0.574775 landard deviation = 0.758139 :andard error = 0.154754 .nlmum = -2.07944 ximum = 1.88707 inge = 3.96651 wer quartile = 1.28093 per quartile = 1.58924 terquartile range = 0.308301 ewness = -4.13299 nd. skewness = -8.26598rtosis = 18.9091 nd. kurtosis = 18.9091 eff. of variation = 58.3324 n = 31.1925

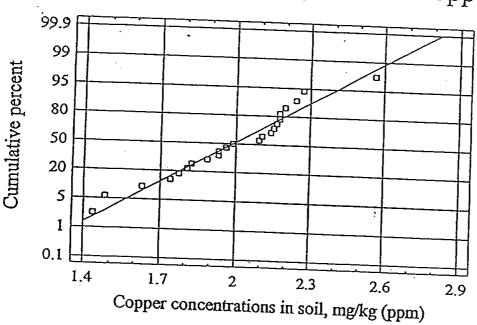
# Lognormal Probability Plot for Cobalt



Summary Statistics for log(Copper)

Count = 23 Average = 1.98556 Median = 1.98787 Geometric mean = 1.96762 Variance = 0.0713494 Standard deviation = 0.267113 Standard error = 0.0556969 Minimum - 1.43508 Maximum = 2.56495 Range = 1.12986 Lower quartile = 1.80829 Upper quartile = 2.17475 Interquartile range = 0.366463 Skewness = -0.263077 Stnd. skewness = -0.515077Kurtosis = 0.18883 Stnd. kurtosis = 0.184854Coeff. of variation = 13.4528 3um = 45.6679

# Lognormal Probability Plot for Copper

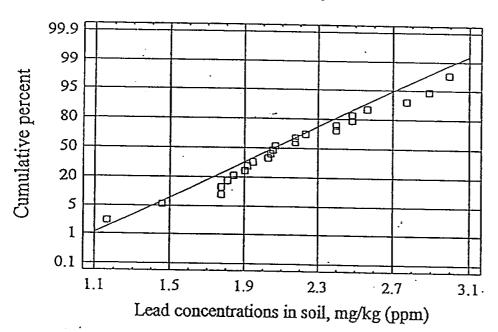


immacy Statistics for log(Lead) 2.13936 :dian = 2.06049 ometric mean = 2.09509 riance = 0.187882 andard deviation = 0.433454 andard error = 0.0884784 nlmum = 1.16315 ximum - 2.99573 nge = 1.83258wer quartile = 1.87133 per quartile = 2.4414 terquartile range = 0.570072 ewness = 0.0350174nd. skewness = 0.0700348 ctosis = 0.200156 nd. kurtosis = 0.200156

of variation = 20.261

n = 51.3446

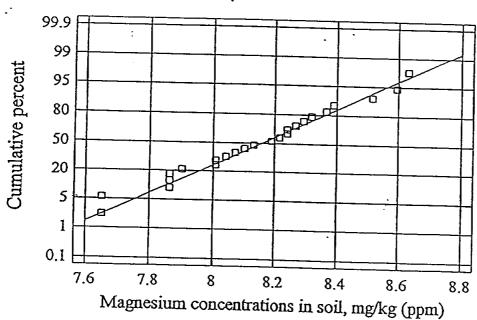
#### Lognormal Probability Plot for Lead



Summary Statistics for log(Magnesium) lount = 24 werage = 0.14232 (edian = 8.16011 iode = eometric mean = 8.13815 'ariance = 0.0706013 tandard deviation = 0.265709 tandard error = 0.0542376inimum = 7.64969 aximum = 8.63052 ange - 0.980829 ower quartile = 7.95369 oper quartile = 8.3064 iterquartile range = 0.352709 cewness = -0.0600481 ind. skewness = -0.120096 irtosis = -0.414246 :nd. kurtosis = -0.414246 peff. of variation = 3.26331

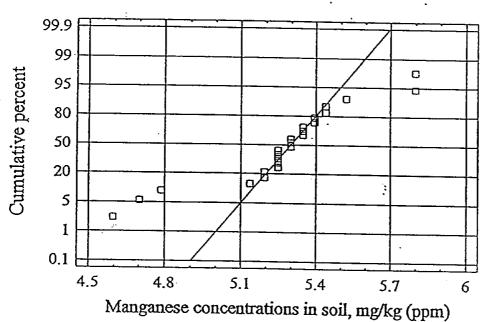
m = 195.416

# Lognormal Probability Plot for Magnesium



ummary Statistics for log(Manganese) **≈** 5.2733 edian = 5.29832 ode = eometric mean = 5.2661 3riance = 0.0771874 tandard deviation = 0.277826 candard error = 0.056711 inimum - 4.59512 3ximum = 5.79909 inge = 1.20397 wer quartile = 5.21999 per quartile = 5.39363 iterquartile range = 0.173637 :ewness = -0.660387nd. skewness = -1.32077rtosis = 1.62566 ind. kurtosis = 1.62566 eff. of variation = 5.26854 m = 126.559

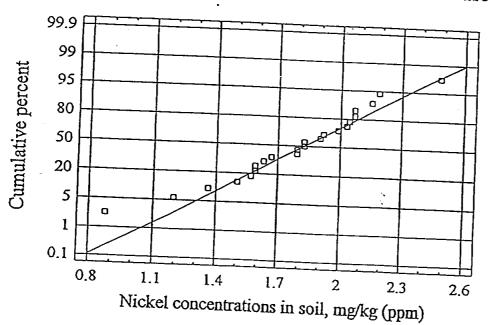
## Lognormal Probability Plot for Manganese



Summary Statistics for log(Nickel)

Count = 23 Average = 1.78451 Median = 1.82455 Mode -Geometric mean = 1.74596 Variance = 0.1246 Standard deviation = 0.352987 Standard error = 0.0736029 Minimum = 0.875469 Maximum - 2.48491 Range = 1.60944 Lower quartile = 1.58924 Upper quartile = 2.04122 Interquartile range = 0.451985 Skewness = -0.609856 Stnd. skewness = -1.19403Kurtosis = 0.992502 Stnd. kurtosis = 0.971605Coeff. of variation = 19.7806 Sum = 41.0438

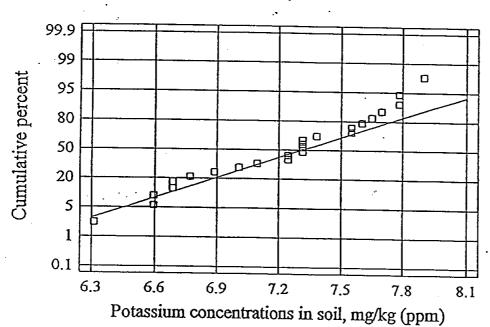
# Lognormal Probability Plot for Nickel



immary Statistics for log(Potassium)

7.21862 7.31322 de = 7.31322 ometric mean = 7.20542 riance = 0.195599 andard deviation = 0.442265 andard error = 0.0902771 nimum = 6.30992ximum = 7.90101nge = 1.59109wer quartile = 6.82802 per quartile = 7.57526 terquartile range = 0.747233 wness = -0.373735 1d. skewness = -0.74747ctosis = -0.83864 nd. kurtosis = -0.83864 of variation = 6.12673 1 = 173.247

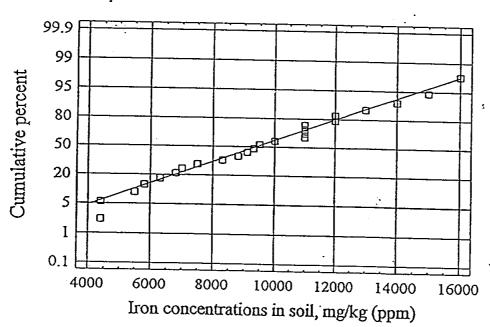
#### Lognormal Probability Plot for Potassium



Summary Statistics for Iron

Count = 24 Average = 9529.17 Median = 9400.0 Mode = 11000.0Geometric mean = 8977.5 Variance = 1.0363E7 Standard deviation = 3219.17 Standard error = 657.109 Minimum = 4400.0 Maximum = 16000.0 Range = 11600.0 Lower quartile = 6900.0 Upper quartile = 11500.0 Interquartile range = 4600.0 Skewness = 0.20025 Stnd. skewness = 0.400499Kurtosis = -0.620589 Stnd. kurtosis = -0.620589 Coeff. of variation = 33.7822 Sum = 228700.0

#### Normal Probability Plot for Iron

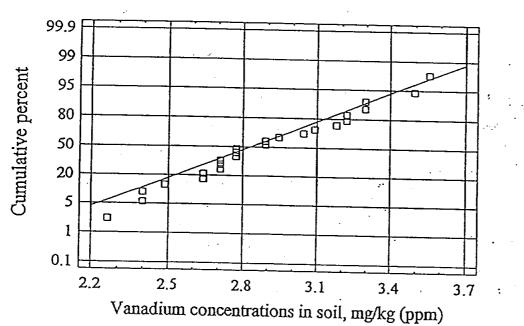


mmary Statistics for log(Vanadium)

= 2.89094
dian = 2.83148
fe =

Description mean = 2.87064
diance = 0.122444
Indard deviation = 0.34992
Indard error = 0.0714271
Indian = 2.26176
dimum = 3.55535
Ige = 1.29358
for quartile = 2.67355
For quartile = 3.19846
for quartile range = 0.524911
Indian = 0.58415
d. skewness = 0.316831
tosis = -0.688491
d. kurtosis = -0.688491
ff. of variation = 12.104
= 69.3826

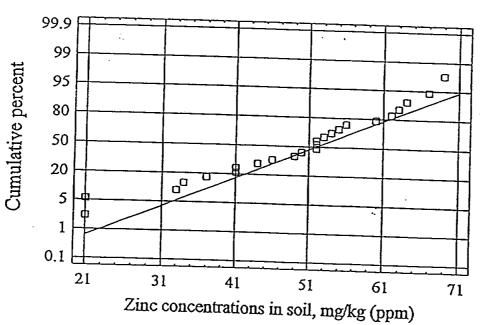
## Lognormal Probability Plot for Vanadium



Summary Statistics for Zinc

Average - 49.0 Median = 52.0 Mode = 52.0 Geometric mean = 46.9434 Variance = 171.478 Standard deviation = 13.095 Standard error = 2.673
Minimum = 21.0
Maximum = 69.0 Range = 48.0 Lower quartile = 41.0 Upper quartile = 58.0 Interquartile range = 17.0 skewness = -0.633044 Stnd. skewness = -1.26609(urtosis = -0.0224531 itnd. kurtosis = -0.0224531 beff. of variation = 26.7244 um = 1176.0

## Normal Probability Plot for Zinc



Results
Soil
Background
Backg
rocal

2 110 ND 0.9 23000 2 130 0.3 1.5 24000 2 150 0.3 1.1 34000 2 150 0.3 1.1 34000 2 210 0.3 2.3 39000 3 150 0.3 1.1 34000 3 150 0.3 1.1 34000				1/	T.				-			<del>_</del>														
Comparison		lercury									S			Q N		2	2	2	QN	2	9	S	2		2	S
Columbia Solution   Colu		yanganese	٧١	200	200	310	210	250	198	210	330	330	190	200	170	180			-	120						
Column   C		muisəngsA	710	2 6	2100	2200	3700	4400	3000	2600	5600	3900	3800	3400	2600	3000	2600			-	├-	٠			_I	5000
2 110 ND 0.8 23000 2 3 6 6800 2 120 ND 1.4 24000 1 95 0.3 1.5 24000 6 6 9 11000		bea	عارد	7	, m		တ	6	8	9	16	11	œ	7	12		Н	9	_	_	ļ.,	1-	+-	+-		1=
2 110 ND 0.9 23000 3 3 6 Coppet 130 ND 1.8 13000 0.3 1.5 24000 0 0.5 1.0 0.9 1		ton	5800	8800	4400	6300	11000	11000	9300	8300	10000	11000	9100	6800	7000	7500	4400	9500	11000	9200	<u> </u>	├	┼		╀	+
200 0.3 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	ŀ	Copper	၂ဖ	1	4	0	6	6	13	7	14	6	80	7	9	9	4	7	7	ည	ω	0	0	┼-	┼	6
200 0.3 5 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2		tisdo	) m	4	က	က	2	ည	4	4		9	4	4	4	4	밁	4	2	က	വ	4	ည	9	2	9
200 0.3 5 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2		тијтол4О	<u>س</u>	2	7	က	ဖ	7	6	4	9	ဖ	19	4	4	4	က	က	9	က	8	7	8	9	7	5
200 0.3 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	on incide	muiɔlsɔ	23000	24000	35000	31000	36000	28000	24000	24000	78000	33000	46000	51000	34000.	39000	54000	00009	43000	82000	42000	35000	43000	-	55000	47000
200 0.3 5 0.		Cadmium	0.9	1.5	0.8	1	8.	1.8	2:3	4.4	1.8	1.7	1.5	-		6.	9.0	9	<u>                                     </u>	6.9	2.3	2.6	2.2	2.4	2.2	2.6
200 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		Beryllium	ND	0.3	g	9	0.4	9.4	0.3	9	0.5	0.5	0.3	6.3	0.3	6.3		-	-							0.4
THE STATE OF THE PROPERTY OF THE PROPERTY OF THE STATE OF		muisea	110	130	110	9	13	35	20	120	210	5	22		-+		+	-		-	-	$\dashv$				290
		Arsenic	2	7	7	7	ည	~	7	7	Θ	6	9	2		+		╬	+	╅	+	9	-		$\dashv$	9
(1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	L	ynomitnA	9	8	4	~	၈	2	7	۽ ام	2		о (	ρ	ا م	1,	٦٥	1	-	7	=		_	_	_	4
		munimulA	2700	4100	2400	3400	4800	0009	4000	2005	2400	2200	4500	2000	200	2000	2600	+-	╅	╁	┽	-		_	+	8600   1
		sample Identifier	Bkg-01-A	Bkg-01-B	Bkg-02-A	Bkg-02-B	BKg-03-A	BKg-03-B	BKg-04-A	DKG-04-B	A-co-by-	BKg-05-B	BKg-Uo-A	BKG-00-B	BKG-07-A	BK9-07-B	84-00-PAB	BYO OO A	0 00 0 A B	0-60-640	A-01-Bvg	BKg-10-B	BKg-11-A	+	-	BKg-12-B

Concentrations in mg/kg Activities in pCi/g Sample Identifier XX-XX-A - surface soil samples Sample Identifier XX-XX-B - subsurface soil samples Local Background Soil Results

				_,_																				
462-muiner							-	- 6	) )					T			T	T	T	T	4		2 0	0.9
182/35/muine	n						0 28		7.02												S	3 6	12	┪—
862-muiner	n						8	200	3	1	T	1	T			T	<u> </u>	1	$\dagger$	-	1			0.6
862 muinotul	ď						< 0.011	0000	200									-			<0.017	<0.00 0.01 8 1.00 0.00 0.00 0.00 0.00 0.0	<0.017	
₽S/96S muinotul	4						<0.00>	<0.008													╁	┦—	+-	1
ասith l	L						<0.010	<0.022													╂—	╀	-	$\vdash$
2 ni Z	50	63	41	53	56	62	55	52	37	34	52	54	21	21	33	67	41	44	52	49	┿	₩	-	69
muibene\	1	18	9.6	1-	19	15	18	16	22	18	16	14	15	15	12	21	24	14	├-	╁╌	-	╌	<del>                                     </del>	33 (
muilledT	Q.	2	2	2	S	2	Ω N	2	Ð	QN	2	2	2	2	2	<u>Q</u>	QN	2	<u> </u>	-	<u> </u>			QN
muiboS	9	2	2	₽	2	480	QN	N N	8	Q	S	420	Ð	380	Q	430	280	640	S	S	_		N N	620
Silver	S	2	Q.	QN N	QN N	S S	-	ND	ND	ΔN	ΔN	QΝ	QN	₽ Z	ΩN	ΠN	QN	QN	QN	QN	QN N	ND	9	밁
muinələ2	QN	QN	QN	ND	ND	QN	Q Z	2	DN	QN	QN	ND	QN	ND		ND	QN	QΝ						S S
Muissstoq	1500	2000	730	1600	1500	1200	1900	1400	2700	1400	1500	800	870	800			_	550		-	2100		-	1900
Nickel	4	ω	7	5	7	တ	12	വ	6	æ	13	9	ည	വ	က	2	8	ည	9	7	7	8	∹	
Sample Identifier	Bkg-01-A	Bkg-01-B	Bkg-02-A	Bkg-02-B	Bkg-03-A	Bkg-03-B	Bkg-04-A	Bkg-04-B	Bkg-05-A	Bkg-05-B	Bkg-06-A	Bkg-06-B	Bkg-07-A	Bkg-07-B	Bkg-08-A	Bkg-08-B	Bkg-09-A	Bkg-09-B	Bkg-10-A	Bkg-10-B	Bkg-11-A	Bkg-11-B	Bkg-12-A	Bkg-12-B
	Nickel Potassium Selenium Silvet Sodium Thallium Vanadium Zinc Intonium 239/24 Sinc Intonium 238	4 Nickel Potassium Selenium Silvet ND Sodium	## Mickel  ## Mickel  ## Mickel  ## Mickel  ## Mickel  ## Mickel  ## Selenium  ## S	4	1	4	9 7 7 8 6 9 4 Nickel 9 7 7 8 6 9 4 Nickel 9 7 7 8 6 9 7 8 8 8 8 8 9 9 9 9 9 9 9 9 9 9 9 9 9	4 8 4 9 7 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	1	9 5 1 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	## 1700 ND	1	## 1	## 13   14   10   10   10   10   10   10   10	A	1	1	A   A   B   B   A   B   B   B   B   B	A   A   A   A   A   A   A   A   A   A	A   A   B   B   B   B   B   B   B   B	A	A	A	A

Concentrations in mg/kg
Activities in pCi/g
Sample Identifier XX-XX-A - surface soil şamples
Sample Identifier XX-XX-B - subsurface soil samples

N	ormal Pa	rame	ters	for Tije	eras	Arrov	/0 l o/	ral Ma	otal Book						
					7	T,	7	Cai IVIE	Tan Back	grour	id Data	3			
7	Į į	}		1	ľε	ΙĒ	1				Se				٦
Statistical	ļ į	۱٤	nic.	Ę	mium	Ę	1 =	ان			Ę	1	dium	1	
Parameter	Aluminum	Antimo	Arse	arium	89	hromium	oba	oppe	Ç	ad	Mangane	Nickel	anadi		1
median	4300	8.5	2	140	15	0	10	U_	<u> </u>	Le	Σg	∣≗	/ar	Zinc	ı
geometric mean	4579.9		3		2	6	4.2	7.3	9400	7.9	200	6.2	17	52	ł
maximum	10000			144	2	5	3.7	7.3	8977.5	8.5	195	6	18		1
minimum		16	6	210	3	10	6.6	13	16000	20	330	12	<del></del>	47	1
	2200	4.4	2	95	_1	2	0.1	4.2	4400	3.2			35	69	l
		9	_3	149	2	5.5	4.2	7.5	9529.2		99	2.4	9.6	21	ı
standard deviation	2095.4	3	2	40.5	1	2.3	1.3			9.3	202	6.3	19	49	l
normal tolerance	2.309	2.3	2	2.33	2				3219.2	4.2	53.6	2.1	6.9	13	
UTL	4927.4	16	<del>  </del>			2.3	2.3	2.3	2.309	2.3	2.31	2.3	2.3	2.3	
				244	3	11	7.3	12	16962	19	326	11	35	70	

Log	inormal F	aram	eter	s for T	ijera	s Arr	oyo L	ocal I	Vietai Bad	karo	und Da	.4-		
Statistical Parameter	Aluminum	Antimony	Arsenic	Barium	Cadmium	Chromium	Cobalt	opper	CO	ad	Manganese	ckei	ınadium	ပ္က
arithmetic average	8.4294	2.2	1	4.97	0	1.6	1.3	δ.	2 4 2 2 =	<u> </u>		ż	\ \ \ \	Zinc
standard deviation	0.4126	0.3	1	0.27	0	0.5			9.1025	2.1	5.27	1.8	2.9	3.8
normal tolerance	2.309	2.3	2	2.33	2		0.8	0.3	0.3631	0.4	0.28	0.4	0.3	0.3
UTL	9:3821	2.9	2	· 5.6	-4	2.3	2.3	2.3	2.309	2.3	2.31	2.3	2.3	2.3
e <sup>UTL</sup>	11874	19				2.7	3.1	2.6	9.941	3.1	5.91	2.6	3.7	4.6
		13	10	271	4	14	21	14	20764	23	370	14	40	9.0

ufficient data for mercury, selenium, silver, and thallium to calculate statistics. All concentrations in mg/kg

# Summary of Background Concentrations for Radionuclides in Soil

Bismuth-212 324 Bismuth-214 340	Number of Detects	Number of Rejected Samples	Distribution Type	Ranga		Geometric Mean	Median	95" Upper Tolerance Limit	45.04039 G * 30
	17	307	Nonconcurrent	(Recal)		(b/C)(b)	(pCVg)	(pCV <sub>Q</sub> )	(pCV9)
	321	ā	Oilleurandino	0.414-2.7	=	1,1055	1.0		27
Cesium-137 802	561	200	Nonparametric	0.27-1.4	321	0.648	0.6		9
(Surface)	t	€ 1	Nonoaramatric	1 000	1			,	5   1
t (B)	ı	ı	Unknown	detection limit	22 52	0.200 <delection limit<="" td=""><td>· 0.2495</td><td>1 (</td><td>0.92</td></delection>	· 0.2495	1 (	0.92
Cobalt-60 321	==	7.7	1.00	(<0.0688)		(<0.0586)	(<0.0686)		(<0.0686)
Lead-210°	ţ		- UNKUOWU	- <detection limit<br="">(&lt;0.0418)</detection>	247	<detection (<0.0418)<="" limit="" td=""><td>cdetection limit</td><td></td><td>«delection limit</td></detection>	cdetection limit		«delection limit
	40	292	Nonparametric	0.3-12.0	46	2 26838	(2)		(<0.0418)
323	233	06	Lognormal	7,1-1,0	2 8	4,4,000	2.835	•	6.8
Lead-214* 249	241	6	Loonormal	- 1 - 00 O	3 3	0.49689	0.5	1.0795	'
Potassium-40 722	720	,	Normal	0.63-1.13	240	0.549	0.56	05.0	'
Radium-224 24	24		i di lina	0.192-31.0	718	15.889	16.4	25.34	'
Radium-226 368	53		Nonparametric	0.43-0.97	25	0.6747	0.655	-	0.558
Radium-228	1	410	Lognormal	0.5-2.09	54	0.713	0.590	76	
1	5	0	Nonparametric	0.45-1.05	22	0,695	0.630		
60	0	0	Unknown		0			•	50.7
1	\$5	65	Nonparametric	0.0321.85	5	0.36.0		•	
136 136	136	0	Lognormal	00,7	:   3	0.5-300	0.2883		0.766
Thorium-234 365	52	330	Lognormal	03071.60	2 3	0.7971	0.810	1.258	,
Tritium	o	0	Unknown	0.0541-0.0	g (	0.7796	0.71	2,89	٠
Uranium-234 4	4	0	olylo Herecook		-  - -	τ	-		,
Uranium-235 95	12	7.5	Simplify of N	0.1-0.0	•	0.897	6.0		1.0
Uranium-238 223	208		incipalatifallic	0.05-0.18	20	0,1198	0.1235	,	0.168
	200	1/	Nonparametric	0.0033-2,065	206	0.506	0.763		=

STATE OF STREET

# Appendix E Summary of Reconnaissance Data Report

The Site 50 reconnaissance study included analysis of eleven soil samples. The sample locations are shown in Figure E-1. Most of the analyses were non-detections. The non-detected parameters, the applied methods, and detection limits are listed in Tables E-1 and E-2. The detected constituents and the range of concentrations are shown in Table E-3. The level of quality assurance/quality control corresponded to EPA procedures and standard commercial standards available in 1987. Five SNL/NM Environmental Program procedures were followed in sample collection:

- 1.1 General Instructions for Field Personnel (revision 0)
- 1.3 Sample Control and Documentation (revision 0)
- 1.4 Sample Containers and Preservation (revision 0)
- 1.6 General Equipment Decontamination (revision 0)
- 5.2 Soil Sampling with a Spade and Scoop (revision 0)

One rinsate blank was collected for the 10 field samples and was analyzed for HSL inorganic constituents. One "duplicate" sample 1 was collected and the relative percent difference between the original sample and the duplicate was calculated and reported for detected constituents.

Table E-1. Non-Detected Constituents (inorganic)

Constituent	Analytical Method	Detection Limit (mg/kg)	Constituent	Analytical Method	Detection Limit (mg/kg)
Silver	HSL Inorganic	1.8 to 2.0	Silver	TCLP Metals	0.5
Beryllium	HSL Inorganic	0.91 to 1.0	Arsenic	TCLP Metals	0.5
Cadmium	HSL Inorganic	0.91 to 1.0	Cadmium	TCLP Metals	0.1
Cerium	HSL Inorganic	18.2 to 20.0	Chromium	TCLP Metals	0.5
Cobalt	HSL Inorganic	9.1 to 10.0	Mercury	TCLP Metals	0.2
Mercury	HSL Inorganic	0.1	Lead	TCLP Metals	0.5
Nickel	HSL Inorganic	7.3 to 8.0	Selenium	TCLP Metals	0.1
Antimony	HSL Inorganic	11.5 to 12.0	Silver	EP Toxicity	0.1 to 0.5
Thallium	HSL Inorganic	1.9 to 2.0	Arsenic	EP Toxicity	0.5
Zirconium	HSL Inorganic	18.2 to 20.1	Cadmium	EP Toxicity	0.5
Chromium	EP Toxicity	0.5	Lead	EP Toxicity	0.5
Mercury	EP Toxicity	0.0002	Selenium	EP Toxicity	0.1

<sup>&</sup>lt;sup>1</sup> The sample was not a true duplicate because it was taken at the same location but at a slightly different depth.

Table E-2. Non-Detected Constituents (organic)

	` ` `	, ,
Constituent	Analytical Method	Detection Limit (mg/kg)
2,4,6-TNT	USATHAMA LW02	0.00192
Gamma hexachlorocyclohexane	EPA 509B	0.00005
Heptachlor	EPA 509B	0.00005
Chlordane	EPA 509B	0.0005
Endrin	EPA 509B	0.0001
Toxaphene	EPA 509B	0.001
Methoxychlor	EPA 509B	0.0005
2,4-Dichlorophenoxyacetic acid	EPA 509B	0.001
2,4,5-Trichlorophenoxypropionic acid	EPA 509B	0.0005
2,4,5-Trichlorophenoxyacetic acid	EPA 509B	0.0005
Phenol	EPA, CLP, GC/MS	0.01
Pyridine	EPA, CLP, GC/MS	0.01
bis (2-chloro-ethyl) ether	EPA, CLP, GC/MS	0.01
1,4-Dichlorobenzene	EPA, CLP, GC/MS	0.01
1,2-Dichlorobenzene	EPA, CLP, GC/MS	0.01
2-Methyl-phenol	EPA, CLP, GC/MS	0.01
3-Methyl-phenol	EPA, CLP, GC/MS	0.01
4-Methyl-phenol	EPA, CLP, GC/MS	0.01
Hexachloroethane	EPA, CLP, GC/MS	0.01

Table E-3. Range of Concentrations (HSL inorganic constituents in mg/kg and Uranium in pCi/g and vg/g)

Constituent	Maximum	Minimum	Constituent	Maximum	Minimum
Aluminum	8750	3570	Magnesium	5620	2770
Arsenic	2.8	1.9 (ND)	Manganese	282	50.1
Barium	286	86.8	Sodium	5880	908 (ND)
Calcium	157000	22500	Nickel	7.9	7.3 (ND)
Chromium	7.7	2.5	Lead	11.2	1.8
Copper	16.3	7.7	Titanium	369	38.3
Iron	8880	3060	Vanadium	25.2	14.1
Potassium	2160	985 (ND)	Zinc	25.3	7.3
U-234	1.0±0.2 pCi/g	0.4±0.2 pCi/g	U-235	0.06±0.06 pCi/g	0.00±0.05 pCi/g
U-238	1.1±0.2 pCi/g	0.4±0.1 pCi/g	Total U	1.4 ug/g	0.91 vg/g

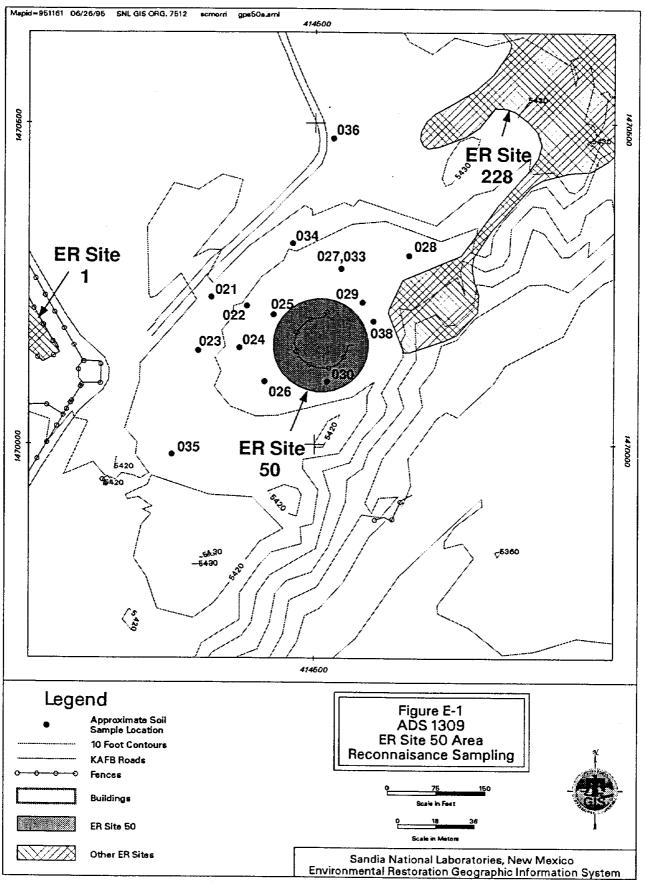


Figure E-1. Old Centrifuge Site 50